

HUMAN RALGDS-LIKE PROTEIN 3

CROSS REFERENCE TO RELATED APPLICATIONS

5 This application claims priority under 35 U.S.C. § 365(c) to international patent application no. PCT/US01/00663, PCT/US01/00664, PCT/US01/00665, PCT/US01/00666, PCT/US01/00667, PCT/US01/00668, PCT/US01/00669 and PCT/US01/00670, all filed January 30, 2001; claims priority under 35 U.S.C. § 120 to 10 commonly owned and copending U.S. application serial no. 09/864,761, filed May 23, 2001; claims priority to United States provisional application serial no: 60/326,105, filed September 28, 2001; the disclosures of which are incorporated herein by reference in their entireties.

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REFERENCE TO SEQUENCE LISTING SUBMITTED ON COMPACT DISC

The present application includes a Sequence Listing filed on a single CD-R disc, provided in duplicate, containing 20 a single file named pto_PB0176.txt, having 33 kilobytes, last modified on January 17, 2002 and recorded January 24, 2002. The Sequence Listing contained in said file on said disc is incorporated herein by reference in its entirety.

25 FIELD OF THE INVENTION

The present invention relates to novel human RalGDS-like protein 3. More specifically, the invention provides isolated nucleic acid molecules encoding human RalGDS-like 30 protein 3, fragments thereof, vectors and host cells comprising isolated nucleic acid molecules encoding human RalGDS-like protein 3, human RalGDS-like protein 3 polypeptides, antibodies, transgenic cells and non-human organisms, and

diagnostic, therapeutic, and investigational methods of using the same.

BACKGROUND OF THE INVENTION

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The Ras superfamily of monomeric GTPases is coupled with receptor and non-receptor tyrosine kinase activation of downstream cytoplasmic and nuclear events. Such events lead to a wide variety of cellular processes, including proliferation, differentiation, nuclear transport, cytoskeleton organization, and vesicular transport. Campbell et al., *Oncogene* 17:1395-1413 (1998), Macara et al., *FASEB J.* 10:625-630 (1996), Hall et al., *Science* 279:509-514 (1998). These GTPases share the ability to cycle between inactive GDP- and active GTP-bound structural states. In the active GTP-bound state, Ras-related proteins interact with a variety of cellular targets to elicit their biological effects. Bourne et al., *Nature* 349:117-127 (1991). The activation of Ras proteins, through the dissociation of GDP and subsequent binding of GTP, is catalyzed by guanine nucleotide exchange factors (GEFs). Quilliam et al., *BioEssays* 17, 395-404 (1995). The return to the inactive GDP-bound state is stimulated by GTPase-activating proteins that promote rapid hydrolysis of GTP. Boguski et al., *Nature* 366, 643-654 (1993).

In searching for the components of Ras signal transduction pathway, multiple Ras effectors have been identified. The best-characterized Ras effectors are the Raf family of serine-threonine kinases including Raf1, B-Raf, and A-Raf. Robinson et al., *Curr. Opin. Cell Biol.* 9:180-186 (1997). Once activated, Raf kinases trigger a cascade of protein kinases that result in the activation of extracellular signal-regulated kinases 1 and 2 (Marshall, *Mol. Reprod. Dev.* 42:493-499 (1995)) which eventually lead to the stimulation of various transcription factors that regulate the expression of

genes involved in proliferation and oncogenesis. Treisman, *Curr. Opin. Cell Biol.* 8:205-215 (1996). In addition to Raf, Ras proteins can also activate other effectors including RalGEF proteins and phosphatidylinositol 3-kinase (PI-3K). Bos et al., 5 *EMBO J.* 17:6776-6782 (1998), Rodriguez-Viciiana et al., *Nature* 370:527-532 (1994). The RalGEF proteins serve as guanine nucleotide exchange factors for the Ras-like GTPase Ral. The activation of PI-3K leads to the activation of both Rho family GTPases and the protein kinase AKT. Franke et al., *Cell* 81:727-10 736 (1995).

To better understand the Ras signaling pathways, Rit - a Ras-related GTPase - was identified. Rit shares about 50% amino acid sequence identity with Ras, and when activated, induces strong growth transformation to NIH3T3 cells. Shao et 15 al., *J. Biol. Chem.* 275:26914-26924 (2000). The Rit-transformed cells proliferate in low serum, form colonies in soft agar, and form tumors in nude mice. Furthermore, Rit stimulates transcription from reporter constructs controlled by minimal promoters containing recognition sites for SRF, NF-B, Elk, and 20 Jun. However, no activation of extracellular signal-regulated kinases, c-Jun N-terminal kinase, or p38, or of PI-3K/Akt/PKB kinases was observed. A combination of biochemical and yeast two-hybrid studies suggest that Rit may interact with only a limited subset of the known Ras-binding proteins, suggesting 25 that Rit might use unique signaling pathways to regulate cellular proliferation and transformation.

In searching for effectors of Rit, a novel RalGEF like mouse protein, RGL3, was identified as a candidate effector for both Ras and Rit. Mouse RGL3 shares 35% amino acid 30 sequence identity with the known RalGEFs. It interacts in a GTP- and effector loop-dependent manner with Rit and Ras through a C-terminal 99-amino acid domain (RA domain). It also exhibits guanine nucleotide exchange activity to the small

GTPase Ral that was stimulated in vivo by the expression of either activated Rit or Ras. Therefore, mouse RGL3 function as an exchange factor for Ral and may serve as a downstream effector for both Rit and Ras.

5 The mouse RGL3 contains three functional domains. The RasGEFN domain is located close to the N-terminal end and is identified in a subset of guanine nucleotide exchange factors for Ras-like small GTPases. The recent crystal structure shows that this domain is alpha-helical and plays a "purely
10 structural role". Boriack-Sjodin et al, *Nature* 394, 337-343 (1998). The RasGEF domain is located at the center of the molecule and may function as the prime site for guanine nucleotide binding and exchange. The RA (Ras association) domain is located at the C-terminal end and is a putative
15 binding site for RasGTP effectors. Gao et al., *J. Biol. Chem.* 10.1074/jbc.M105760200 (2001).

Mouse RGL3 is the latest member of RalGEF family proteins identified as an effector of Ras proteins. Ehrhardt et al, *Oncogene* 20(2):188-97 (2001), Shao et al., *J. Biol. Chem.* 20 275:26914-26924 (2000). It is unknown whether these RalGEFs are functionally redundant or if they have distinct physiological roles. However, the fact that the mouse RGL3 is only 35% identical to other RalGEF family proteins at the amino acid level suggests that mouse RGL3 may play a unique role in
25 mediating cellular signaling. The fact that mouse RGL3 interacts with both Ras and Rit also supports this notion. Through mediation of cellular signaling, mouse RGL3 likely regulates cellular proliferation and transformation. This would argue strongly that similar to the Ras and Rit proteins
30 upstream in the signaling pathway, mouse RGL3 is also an oncogene.

Given a likely role of mouse RGL3 in signal transduction and oncogenesis, it has potential therapeutic as

well as diagnostic roles in a variety of human tumors. There is a need to identify and to characterize human forms of the RGL3 protein.

5 SUMMARY OF THE INVENTION

The present invention solves these and other needs in the art by providing isolated nucleic acids that encode human RalGDS-like protein 3 (RGL3), and fragments thereof.

10 In other aspects, the invention provides vectors for propagating and expressing the nucleic acids of the present invention, host cells comprising the nucleic acids and vectors of the present invention, proteins, protein fragments, and protein fusions of the RGL3, and antibodies thereto.

15 The invention further provides pharmaceutical formulations of the nucleic acids, proteins, and antibodies of the present invention.

20 In other aspects, the invention provides transgenic cells and non-human organisms comprising RGL3 nucleic acids, and transgenic cells and non-human organisms with targeted disruption of the endogenous orthologue of the RGL3.

25 The invention additionally provides diagnostic, investigational, and therapeutic methods based on the RGL3 nucleic acids, proteins, and antibodies of the present invention.

BRIEF DESCRIPTION OF THE DRAWINGS

30 The above and other objects and advantages of the present invention will be apparent upon consideration of the following detailed description taken in conjunction with the accompanying drawings, in which like characters refer to like parts throughout, and in which:

FIG. 1 (A) schematizes the protein domain structure of RGL3, FIG. 1 (B) shows the alignment of RasGEFN domain of RGL3 with that of other proteins, FIG. 1 (C) shows the alignment of RasGEF domain of RGL3 with that of other proteins,
5 and FIG. 1 (D) shows the alignment of RA domain of RGL3 with that of other proteins;

FIG. 2 is a map showing the genomic structure of RGL3 encoded at chromosome 19p13.2;

10 FIG. 3 presents the nucleotide and predicted amino acid sequences of RGL3; and

FIG. 4 presents the expression analysis of RGL3 by RT-PCR.

DETAILED DESCRIPTION OF THE INVENTION

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Mining the sequence of the human genome for novel human genes, the present inventors have identified RGL3, a guanine nucleotide exchange factor for the small GTPase Ral and a downstream effector for both Rit and Ras. Through the
20 mediation of cellular signaling, RGL3 likely regulates cellular proliferation and transformation.

As schematized in FIG. 1, the newly isolated gene product shares certain protein domains and an overall structural organization with mouse RGL3 and other RasGEF molecules. The shared structural features strongly imply that RGL3 plays a role similar to that of mouse RGL3 as a guanine nucleotide exchange factor for the small GTPase Ral and a downstream effector for both Rit and Ras. Through the mediation of cellular signaling, RGL3 likely regulates cellular
25 proliferation and transformation.

Like mouse RGL3 and other RasGEF molecules, RGL3 contains three functional domains. A RasGEFN domain is located close to the N-terminal end (amino acid sequence 64 - 198), a

RasGEF domain is found at the center of the molecule (amino acid sequence 243 - 506) and a RA (Ras association) domain is found at the C-terminal end (613 - 699).

- <http://www.ncbi.nlm.nih.gov/Structure/cdd/wrpsb.cgi>. The
5 RasGEFN domain is identified in a subset of guanine nucleotide exchange factor for Ras-like small GTPases. Recent crystal structure of the RasGEFN domain shows that this domain is alpha-helical and plays a "purely structural role". Boriack-Sjodin et al, *Nature* 394, 337-343 (1998). The RasGEF domain may
10 function as the prime site for guanine nucleotide binding and exchange and the RA is a putative binding site for RasGTP effectors. Shao et al., *J. Biol. Chem.* 275:26914-26924 (2000).

Other signatures of the newly isolated RGL3 protein were identified by searching the PROSITE database,
15 (<http://www.expasy.ch/tools/scnpsit1.html>). These include one N-glycosylation sites (339-342), three cAMP- and cGMP-dependent protein kinase phosphorylation site (374-377, 517-520, 523-526), fourteen protein kinase C phosphorylation sites (30-32, 36-38, 63-65, 99-101, 170-172, 277-279, 290-292, 342-344, 372-374, 388-390, 495-497, 512-514, 559-561, and 591-593), nine casein kinase II phosphorylation sites (40-43, 221-224, 247-250, 256-259, 277-280, 388-391, 402-405, 540-543, and 635-638), and thirteen N-myristoylation sites (24-29, 169-174, 181-186, 273-278, 283-288, 286-291, 303-308, 307-312, 410-415, 425-430, 25 554-559, 568-573, and 700-705).

FIG. 2 shows the genomic organization of RGL3.

At the top is shown the two bacterial artificial chromosomes (BACs), with GenBank accession numbers (AC008481.8 and AC024575.5), that span the RGL3 locus. The genome-derived
30 single-exon probe first used to demonstrate expression from this locus is shown below the BACs and labeled "500". The 500 bp probe includes sequence drawn solely from exon five.

As shown in FIG. 2, RGL3 encodes a protein of 710 amino acids and is comprised of exons 1 - 19. The predicted molecular weight of RGL3, prior to any post-translational modification, is 78.1 kD.

5 As further discussed in the examples herein, expression of RGL3 was assessed using hybridization to genome-derived single exon microarrays. Microarray analysis of exon five showed expression in all tissues tested, including adrenal, adult liver, bone marrow, brain, fetal liver, heart, 10 kidney, lung, placenta and prostate, as well as a cell line, HeLa. This was confirmed by RT-PCR analysis.

As more fully described below, the present invention provides isolated nucleic acids that encode RGL3 and fragments thereof. The invention further provides vectors for 15 propagation and expression of the nucleic acids of the present invention, host cells comprising the nucleic acids and vectors of the present invention, proteins, protein fragments, and protein fusions of the present invention, and antibodies specific for all or any one of the isoforms. The invention 20 provides pharmaceutical formulations of the nucleic acids, proteins, and antibodies of the present invention. The invention further provides transgenic cells and non-human organisms comprising human RGL3 nucleic acids, and transgenic cells and non-human organisms with targeted disruption of the 25 endogenous orthologue of the human RGL3. The invention additionally provides diagnostic, investigational, and therapeutic methods based on the RGL3 nucleic acids, proteins, and antibodies of the present invention.

DEFINITIONS

Unless defined otherwise, all technical and scientific terms used herein have the meaning commonly understood by one of ordinary skill in the art to which this invention belongs.

As used herein, "**nucleic acid**" (synonymously, "**polynucleotide**") includes polynucleotides having natural nucleotides in native 5'-3' phosphodiester linkage - e.g., DNA or RNA - as well as polynucleotides that have nonnatural nucleotide analogues, nonnative internucleoside bonds, or both, so long as the nonnatural polynucleotide is capable of sequence-discriminating basepairing under experimentally desired conditions. Unless otherwise specified, the term "nucleic acid" includes any topological conformation; the term thus explicitly comprehends single-stranded, double-stranded, partially duplexed, triplexed, hairpinned, circular, and padlocked conformations.

As used herein, an "**isolated nucleic acid**" is a nucleic acid molecule that exists in a physical form that is nonidentical to any nucleic acid molecule of identical sequence as found in nature; "isolated" does not require, although it does not prohibit, that the nucleic acid so described has itself been physically removed from its native environment.

For example, a nucleic acid can be said to be "isolated" when it includes nucleotides and/or internucleoside bonds not found in nature. When instead composed of natural nucleosides in phosphodiester linkage, a nucleic acid can be said to be "isolated" when it exists at a purity not found in nature, where purity can be adjudged with respect to the presence of nucleic acids of other sequence, with respect to the presence of proteins, with respect to the presence of lipids, or with respect the presence of any other component of

a biological cell, or when the nucleic acid lacks sequence that flanks an otherwise identical sequence in an organism's genome, or when the nucleic acid possesses sequence not identically present in nature.

5 As so defined, "isolated nucleic acid" includes nucleic acids integrated into a host cell chromosome at a heterologous site, recombinant fusions of a native fragment to a heterologous sequence, recombinant vectors present as episomes or as integrated into a host cell chromosome.

10 As used herein, an isolated nucleic acid "encodes" a reference polypeptide when at least a portion of the nucleic acid, or its complement, can be directly translated to provide the amino acid sequence of the reference polypeptide, or when the isolated nucleic acid can be used, alone or as part of an 15 expression vector, to express the reference polypeptide *in vitro*, in a prokaryotic host cell, or in a eukaryotic host cell.

As used herein, the term "**exon**" refers to a nucleic acid sequence found in genomic DNA that is bioinformatically 20 predicted and/or experimentally confirmed to contribute contiguous sequence to a mature mRNA transcript.

As used herein, the phrase "**open reading frame**" and the equivalent acronym "**ORF**" refer to that portion of a transcript-derived nucleic acid that can be translated in its 25 entirety into a sequence of contiguous amino acids. As so defined, an ORF has length, measured in nucleotides, exactly divisible by 3. As so defined, an ORF need not encode the entirety of a natural protein.

As used herein, the phrase "**ORF-encoded peptide**" 30 refers to the predicted or actual translation of an ORF.

As used herein, the phrase "**degenerate variant**" of a reference nucleic acid sequence intends all nucleic acid sequences that can be directly translated, using the standard

genetic code, to provide an amino acid sequence identical to that translated from the reference nucleic acid sequence.

As used herein, the term "**microarray**" and the equivalent phrase "**nucleic acid microarray**" refer to a 5 substrate-bound collection of plural nucleic acids, hybridization to each of the plurality of bound nucleic acids being separately detectable. The substrate can be solid or porous, planar or non-planar, unitary or distributed.

As so defined, the term "microarray" and phrase 10 "nucleic acid microarray" include all the devices so called in Schena (ed.), DNA Microarrays: A Practical Approach (Practical Approach Series), Oxford University Press (1999) (ISBN: 0199637768); *Nature Genet.* 21(1)(suppl):1 - 60 (1999); and Schena (ed.), Microarray Biochip: Tools and Technology, Eaton 15 Publishing Company/BioTechniques Books Division (2000) (ISBN: 1881299376), the disclosures of which are incorporated herein by reference in their entireties.

As so defined, the term "microarray" and phrase "nucleic acid microarray" also include substrate-bound 20 collections of plural nucleic acids in which the plurality of nucleic acids are distributably disposed on a plurality of beads, rather than on a unitary planar substrate, as is described, *inter alia*, in Brenner et al., *Proc. Natl. Acad. Sci. USA* 97(4):166501670 (2000), the disclosure of which is 25 incorporated herein by reference in its entirety; in such case, the term "microarray" and phrase "nucleic acid microarray" refer to the plurality of beads in aggregate.

As used herein with respect to solution phase hybridization, the term "**probe**", or equivalently, "**nucleic acid probe**" or "**hybridization probe**", refers to an isolated nucleic 30 acid of known sequence that is, or is intended to be, detectably labeled. As used herein with respect to a nucleic acid microarray, the term "probe" (or equivalently "nucleic

acid probe" or "hybridization probe") refers to the isolated nucleic acid that is, or is intended to be, bound to the substrate. In either such context, the term "target" refers to nucleic acid intended to be bound to probe by sequence complementarity.

As used herein, the expression "**probe comprising SEQ ID NO:X**", and variants thereof, intends a nucleic acid probe, at least a portion of which probe has either (i) the sequence directly as given in the referenced SEQ ID NO:X, or (ii) a sequence complementary to the sequence as given in the referenced SEQ ID NO:X, the choice as between sequence directly as given and complement thereof dictated by the requirement that the probe be complementary to the desired target.

As used herein, the phrases "**expression of a probe**" and "**expression of an isolated nucleic acid**" and their linguistic equivalents intend that the probe or, (respectively, the isolated nucleic acid), or a probe (or, respectively, isolated nucleic acid) complementary in sequence thereto, can hybridize detectably under high stringency conditions to a sample of nucleic acids that derive from mRNA transcripts from a given source. For example, and by way of illustration only, expression of a probe in "liver" means that the probe can hybridize detectably under high stringency conditions to a sample of nucleic acids that derive from mRNA obtained from liver.

As used herein, "**a single exon probe**" comprises at least part of an exon ("reference exon") and can hybridize detectably under high stringency conditions to transcript-derived nucleic acids that include the reference exon. The single exon probe will not, however, hybridize detectably under high stringency conditions to nucleic acids that lack the reference exon and that consist of one or more

exons that are found adjacent to the reference exon in the genome.

For purposes herein, "**high stringency conditions**" are defined for solution phase hybridization as aqueous hybridization (i.e., free of formamide) in 6X SSC (where 20X SSC contains 3.0 M NaCl and 0.3 M sodium citrate), 1% SDS at 65°C for at least 8 hours, followed by one or more washes in 0.2X SSC, 0.1% SDS at 65°C. "**Moderate stringency conditions**" are defined for solution phase hybridization as aqueous hybridization (i.e., free of formamide) in 6X SSC, 1% SDS at 65°C for at least 8 hours, followed by one or more washes in 2x SSC, 0.1% SDS at room temperature.

For microarray-based hybridization, standard "high stringency conditions" are defined as hybridization in 50% formamide, 5X SSC, 0.2 µg/µl poly(dA), 0.2 µg/µl human cot1 DNA, and 0.5% SDS, in a humid oven at 42°C overnight, followed by successive washes of the microarray in 1X SSC, 0.2% SDS at 55°C for 5 minutes, and then 0.1X SSC, 0.2% SDS, at 55°C for 20 minutes. For microarray-based hybridization, "moderate stringency conditions", suitable for cross-hybridization to mRNA encoding structurally- and functionally-related proteins, are defined to be the same as those for high stringency conditions but with reduction in temperature for hybridization and washing to room temperature (approximately 25°C).

As used herein, the terms "**protein**", "**polypeptide**", and "**peptide**" are used interchangeably to refer to a naturally-occurring or synthetic polymer of amino acid monomers (residues), irrespective of length, where amino acid monomer here includes naturally-occurring amino acids, naturally-occurring amino acid structural variants, and synthetic non-naturally occurring analogs that are capable of participating in peptide bonds. The terms "**protein**", "**polypeptide**", and

"peptide" explicitly permits of post-translational and post-synthetic modifications, such as glycosylation.

The term "**oligopeptide**" herein denotes a protein, polypeptide, or peptide having 25 or fewer monomeric subunits.

5 The phrases "**isolated protein**", "**isolated polypeptide**", "**isolated peptide**" and "**isolated oligopeptide**" refer to a protein (or respectively to a polypeptide, peptide, or oligopeptide) that is nonidentical to any protein molecule of identical amino acid sequence as found in nature; "isolated" 10 does not require, although it does not prohibit, that the protein so described has itself been physically removed from its native environment.

For example, a protein can be said to be "isolated" when it includes amino acid analogues or derivatives not found 15 in nature, or includes linkages other than standard peptide bonds.

When instead composed entirely of natural amino acids linked by peptide bonds, a protein can be said to be "isolated" when it exists at a purity not found in nature – where purity 20 can be adjudged with respect to the presence of proteins of other sequence, with respect to the presence of non-protein compounds, such as nucleic acids, lipids, or other components of a biological cell, or when it exists in a composition not found in nature, such as in a host cell that does not naturally 25 express that protein.

A "**purified protein**" (equally, a purified polypeptide, peptide, or oligopeptide) is an isolated protein, as above described, present at a concentration of at least 95%, as measured on a weight basis with respect to total protein in 30 a composition. A "**substantially purified protein**" (equally, a substantially purified polypeptide, peptide, or oligopeptide) is an isolated protein, as above described, present at a

concentration of at least 70%, as measured on a weight basis with respect to total protein in a composition.

As used herein, the phrase "**protein isoforms**" refers to a plurality of proteins having nonidentical primary amino acid sequence but that share amino acid sequence encoded by at least one common exon.

As used herein, the phrase "**alternative splicing**" and its linguistic equivalents includes all types of RNA processing that lead to expression of plural protein isoforms from a single gene; accordingly, the phrase "**splice variant(s)**" and its linguistic equivalents embraces mRNAs transcribed from a given gene that, however processed, collectively encode plural protein isoforms. For example, and by way of illustration only, splice variants can include exon insertions, exon extensions, exon truncations, exon deletions, alternatives in the 5' untranslated region ("5' UT") and alternatives in the 3' untranslated region ("3' UT"). Such 3' alternatives include, for example, differences in the site of RNA transcript cleavage and site of poly(A) addition. See, e.g., Gautheret et al., *Genome Res.* 8:524-530 (1998).

As used herein, "**orthologues**" are separate occurrences of the same gene in multiple species. The separate occurrences have similar, albeit nonidentical, amino acid sequences, the degree of sequence similarity depending, in part, upon the evolutionary distance of the species from a common ancestor having the same gene.

As used herein, the term "**paralogues**" indicates separate occurrences of a gene in one species. The separate occurrences have similar, albeit nonidentical, amino acid sequences, the degree of sequence similarity depending, in part, upon the evolutionary distance from the gene duplication event giving rise to the separate occurrences.

As used herein, the term "homologues" is generic to "orthologues" and "paralogues".

As used herein, the term "antibody" refers to a polypeptide, at least a portion of which is encoded by at least one immunoglobulin gene, or fragment thereof, and that can bind specifically to a desired target molecule. The term includes naturally-occurring forms, as well as fragments and derivatives.

Fragments within the scope of the term "antibody" include those produced by digestion with various proteases, those produced by chemical cleavage and/or chemical dissociation, and those produced recombinantly, so long as the fragment remains capable of specific binding to a target molecule. Among such fragments are Fab, Fab', Fv, F(ab)'₂, and single chain Fv (scFv) fragments.

Derivatives within the scope of the term include antibodies (or fragments thereof) that have been modified in sequence, but remain capable of specific binding to a target molecule, including: interspecies chimeric and humanized antibodies; antibody fusions; heteromeric antibody complexes and antibody fusions, such as diabodies (bispecific antibodies), single-chain diabodies, and intrabodies (see, e.g., Marasco (ed.), Intracellular Antibodies: Research and Disease Applications, Springer-Verlag New York, Inc. (1998) (ISBN: 3540641513), the disclosure of which is incorporated herein by reference in its entirety).

As used herein, antibodies can be produced by any known technique, including harvest from cell culture of native B lymphocytes, harvest from culture of hybridomas, recombinant expression systems, and phage display.

As used herein, "antigen" refers to a ligand that can be bound by an antibody; an antigen need not itself be

immunogenic. The portions of the antigen that make contact with the antibody are denominated "**epitopes**".

"**Specific binding**" refers to the ability of two molecular species concurrently present in a heterogeneous sample to bind to one another in preference to binding to other molecular species in the sample. Typically, a specific binding interaction will discriminate over adventitious binding interactions in the reaction by at least two-fold, more typically by at least 10-fold, often at least 100-fold; when used to detect analyte, specific binding is sufficiently discriminatory when determinative of the presence of the analyte in a heterogeneous (inhomogeneous) sample. Typically, the affinity or avidity of a specific binding reaction is least about 10^{-7} M, with specific binding reactions of greater specificity typically having affinity or avidity of at least 10^{-8} M to at least about 10^{-9} M.

As used herein, "**molecular binding partners**" – and equivalently, "**specific binding partners**" – refer to pairs of molecules, typically pairs of biomolecules, that exhibit specific binding. Nonlimiting examples are receptor and ligand, antibody and antigen, and biotin to any of avidin, streptavidin, neutrAvidin and captAvidin.

The term "**antisense**", as used herein, refers to a nucleic acid molecule sufficiently complementary in sequence, and sufficiently long in that complementary sequence, as to hybridize under intracellular conditions to (i) a target mRNA transcript or (ii) the genomic DNA strand complementary to that transcribed to produce the target mRNA transcript.

The term "**portion**", as used with respect to nucleic acids, proteins, and antibodies, is synonymous with "fragment".

NUCLEIC ACID MOLECULES

In a first aspect, the invention provides isolated nucleic acids that encode RGL3, variants having at least 65% sequence identity thereto, degenerate variants thereof, variants that encode RGL3 proteins having conservative or moderately conservative substitutions, cross-hybridizing nucleic acids, and fragments thereof.

FIG. 3 presents the nucleotide sequence of the RGL3 cDNA clone, with predicted amino acid translation; the sequences are further presented in the Sequence Listing, incorporated herein by reference in its entirety, in SEQ ID NOS: 1 (full length nucleotide sequence of human RGL3 cDNA) and 3 (full length amino acid coding sequence of human RGL3).

Unless otherwise indicated, each nucleotide sequence is set forth herein as a sequence of deoxyribonucleotides. It is intended, however, that the given sequence be interpreted as would be appropriate to the polynucleotide composition: for example, if the isolated nucleic acid is composed of RNA, the given sequence intends ribonucleotides, with uridine substituted for thymidine.

Unless otherwise indicated, nucleotide sequences of the isolated nucleic acids of the present invention were determined by sequencing a DNA molecule that had resulted, directly or indirectly, from at least one enzymatic polymerization reaction (e.g., reverse transcription and/or polymerase chain reaction) using an automated sequencer (such as the MegaBACE™ 1000, Amersham Biosciences, Sunnyvale, CA, USA), or by reliance upon such sequence or upon genomic sequence prior-accessioned into a public database. Unless otherwise indicated, all amino acid sequences of the polypeptides of the present invention were predicted by translation from the nucleic acid sequences so determined.

As a consequence, any nucleic acid sequence presented herein may contain errors introduced by erroneous incorporation of nucleotides during polymerization, by erroneous base calling by the automated sequencer (although such sequencing errors have been minimized for the nucleic acids directly determined herein, unless otherwise indicated, by the sequencing of each of the complementary strands of a duplex DNA), or by similar errors accessioned into the public database. Such errors can readily be identified and corrected by resequencing of the genomic locus using standard techniques.

Single nucleotide polymorphisms (SNPs) occur frequently in eukaryotic genomes - more than 1.4 million SNPs have already identified in the human genome, International Human Genome Sequencing Consortium, *Nature* 409:860 - 921 (2001) - and the sequence determined from one individual of a species may differ from other allelic forms present within the population. Additionally, small deletions and insertions, rather than single nucleotide polymorphisms, are not uncommon in the general population, and often do not alter the function of the protein.

Accordingly, it is an aspect of the present invention to provide nucleic acids not only identical in sequence to those described with particularity herein, but also to provide isolated nucleic acids at least about 65% identical in sequence to those described with particularity herein, typically at least about 70%, 75%, 80%, 85%, or 90% identical in sequence to those described with particularity herein, usefully at least about 91%, 92%, 93%, 94%, or 95% identical in sequence to those described with particularity herein, usefully at least about 96%, 97%, 98%, or 99% identical in sequence to those described with particularity herein, and, most conservatively, at least about 99.5%, 99.6%, 99.7%, 99.8% and 99.9% identical in sequence to those described with particularity herein. These

sequence variants can be naturally occurring or can result from human intervention, as by random or directed mutagenesis.

For purposes herein, percent identity of two nucleic acid sequences is determined using the procedure of Tatiana et al., "Blast 2 sequences - a new tool for comparing protein and nucleotide sequences", *FEMS Microbiol Lett.* 174:247-250 (1999), which procedure is effectuated by the computer program BLAST 2 SEQUENCES, available online at

<http://www.ncbi.nlm.nih.gov/blast/bl2seq/bl2.html>.

- 10 To assess percent identity of nucleic acids, the BLASTN module of BLAST 2 SEQUENCES is used with default values of (i) reward for a match: 1; (ii) penalty for a mismatch: -2; (iii) open gap 5 and extension gap 2 penalties; (iv) gap X_dropoff 50 expect 10 word size 11 filter, and both sequences are entered in their 15 entireties.

As is well known, the genetic code is degenerate, with each amino acid except methionine translated from a plurality of codons, thus permitting a plurality of nucleic acids of disparate sequence to encode the identical protein.

- 20 As is also well known, codon choice for optimal expression varies from species to species. The isolated nucleic acids of the present invention being useful for expression of RGL3 proteins and protein fragments, it is, therefore, another aspect of the present invention to provide isolated nucleic 25 acids that encode RGL3 proteins and portions thereof not only identical in sequence to those described with particularity herein, but degenerate variants thereof as well.

- As is also well known, amino acid substitutions occur frequently among natural allelic variants, with conservative 30 substitutions often occasioning only *de minimis* change in protein function.

Accordingly, it is an aspect of the present invention to provide nucleic acids not only identical in sequence to

those described with particularity herein, but also to provide isolated nucleic acids that encode RGL3, and portions thereof, having conservative amino acid substitutions, and also to provide isolated nucleic acids that encode RGL3, and portions thereof, having moderately conservative amino acid substitutions.

Although there are a variety of metrics for calling conservative amino acid substitutions, based primarily on either observed changes among evolutionarily related proteins or on predicted chemical similarity, for purposes herein a conservative replacement is any change having a positive value in the PAM250 log-likelihood matrix reproduced herein below (see Gonnet et al., *Science* 256(5062):1443-5 (1992)):

	A	R	N	D	C	Q	E	G	H	I	L	K	M	F	P	S	T	W	Y	V
15	2	-1	0	0	0	0	0	0	-1	-1	-1	0	-1	-2	0	1	1	-4	-2	0
	A	R	N	D	C	Q	E	G	H	I	L	K	M	F	P	S	T	W	Y	V
20	-1	5	0	0	-2	2	0	-1	1	-2	-2	3	-2	-3	-1	0	0	-2	-2	-2
	A	R	N	D	C	Q	E	G	H	I	L	K	M	F	P	S	T	W	Y	V
25	0	0	4	2	-2	1	1	0	1	-3	-3	1	-2	-3	-1	1	0	-4	-1	-2
	A	R	N	D	C	Q	E	G	H	I	L	K	M	F	P	S	T	W	Y	V
30	0	0	2	5	-3	1	3	0	0	-4	-4	0	-3	-4	-1	0	0	-5	-3	-3
	A	R	N	D	C	Q	E	G	H	I	L	K	M	F	P	S	T	W	Y	V
35	0	-2	-2	-3	12	-2	-3	-2	-1	-1	-2	-3	-1	-1	-3	0	0	-1	0	0
	A	R	N	D	C	Q	E	G	H	I	L	K	M	F	P	S	T	W	Y	V
40	0	2	1	1	-2	3	2	-1	1	-2	-2	2	-1	-3	0	0	0	-3	-2	-2
	A	R	N	D	C	Q	E	G	H	I	L	K	M	F	P	S	T	W	Y	V
45	0	0	1	3	-3	2	4	-1	0	-3	-3	1	-2	-4	0	0	0	-4	-3	-2
	A	R	N	D	C	Q	E	G	H	I	L	K	M	F	P	S	T	W	Y	V
50	0	-1	0	0	-2	-1	-1	7	-1	-4	-4	-1	-4	-5	-2	0	-1	-4	-4	-3
	A	R	N	D	C	Q	E	G	H	I	L	K	M	F	P	S	T	W	Y	V
55	-1	1	1	0	-1	1	0	-1	6	-2	-2	1	-1	0	-1	0	0	-1	2	-2
	A	R	N	D	C	Q	E	G	H	I	L	K	M	F	P	S	T	W	Y	V
60	-1	-2	-3	-4	-1	-2	-3	-4	-2	4	3	-2	2	1	-3	-2	-1	-2	-1	3
	A	R	N	D	C	Q	E	G	H	I	L	K	M	F	P	S	T	W	Y	V
65	-1	-2	-3	-4	-2	-2	-3	-4	-2	3	4	-2	3	2	-2	-2	-1	-1	0	2
	A	R	N	D	C	Q	E	G	H	I	L	K	M	F	P	S	T	W	Y	V
70	0	3	1	0	-3	2	1	-1	1	-2	-2	3	-1	-3	-1	0	0	-4	-2	-2
	A	R	N	D	C	Q	E	G	H	I	L	K	M	F	P	S	T	W	Y	V
75	-1	-2	-2	-3	-1	-1	-2	-4	-1	2	3	-1	4	2	-2	-1	-1	-1	0	2
	A	R	N	D	C	Q	E	G	H	I	L	K	M	F	P	S	T	W	Y	V
80	-2	-3	-3	-4	-1	-3	-4	-5	0	1	2	-3	2	7	-4	-3	-2	4	5	0
	A	R	N	D	C	Q	E	G	H	I	L	K	M	F	P	S	T	W	Y	V
85	0	-1	-1	-1	-3	0	0	-2	-1	-3	-2	-1	-2	-4	8	0	0	-5	-3	-2
	A	R	N	D	C	Q	E	G	H	I	L	K	M	F	P	S	T	W	Y	V
90	1	0	1	0	0	0	0	0	0	-2	-2	0	-1	-3	0	2	2	-3	-2	-1
	A	R	N	D	C	Q	E	G	H	I	L	K	M	F	P	S	T	W	Y	V
95	1	0	0	0	0	0	0	-1	0	-1	-1	0	-1	-2	0	2	2	-4	-2	0
	A	R	N	D	C	Q	E	G	H	I	L	K	M	F	P	S	T	W	Y	V
100	-4	-2	-4	-5	-1	-3	-4	-4	-1	-2	-1	-4	-1	4	-5	-3	-4	14	4	-3
	A	R	N	D	C	Q	E	G	H	I	L	K	M	F	P	S	T	W	Y	V
105	-2	-2	-1	-3	0	-2	-3	-4	2	-1	0	-2	0	5	-3	-2	-2	4	8	-1
	A	R	N	D	C	Q	E	G	H	I	L	K	M	F	P	S	T	W	Y	V
110	0	-2	-2	-3	0	-2	-2	-3	-2	3	2	-2	2	0	-2	-1	0	-3	-1	3

For purposes herein, a "moderately conservative" replacement is any change having a nonnegative value in the PAM250 log-likelihood matrix reproduced herein above.

As is also well known in the art, relatedness of
5 nucleic acids can also be characterized using a functional test, the ability of the two nucleic acids to base-pair to one another at defined hybridization stringencies.

It is, therefore, another aspect of the invention to provide isolated nucleic acids not only identical in sequence
10 to those described with particularity herein, but also to provide isolated nucleic acids ("cross-hybridizing nucleic acids") that hybridize under high stringency conditions (as defined herein below) to all or to a portion of various of the isolated RGL3 nucleic acids of the present invention
15 ("reference nucleic acids"), as well as cross-hybridizing nucleic acids that hybridize under moderate stringency conditions to all or to a portion of various of the isolated RGL3 nucleic acids of the present invention.

Such cross-hybridizing nucleic acids are useful,
20 *inter alia*, as probes for, and to drive expression of, proteins related to the proteins of the present invention as alternative isoforms, homologues, paralogues, and orthologues. Particularly useful orthologues are those from other primate species, such as chimpanzee, rhesus macaque, monkey, baboon,
25 orangutan, and gorilla; from rodents, such as rats, mice, guinea pigs; from lagomorphs, such as rabbits; and from domestic livestock, such as cow, pig, sheep, horse, goat and chicken.

For purposes herein, high stringency conditions are
30 defined as aqueous hybridization (*i.e.*, free of formamide) in 6X SSC (where 20X SSC contains 3.0 M NaCl and 0.3 M sodium citrate), 1% SDS at 65°C for at least 8 hours, followed by one or more washes in 0.2X SSC, 0.1% SDS at 65°C. For purposes

herein, moderate stringency conditions are defined as aqueous hybridization (i.e., free of formamide) in 6X SSC, 1% SDS at 65°C for at least 8 hours, followed by one or more washes in 2x SSC, 0.1% SDS at room temperature.

5 The hybridizing portion of the reference nucleic acid is typically at least 15 nucleotides in length, often at least 17 nucleotides in length. Often, however, the hybridizing portion of the reference nucleic acid is at least 20 nucleotides in length, 25 nucleotides in length, and even 30
10 nucleotides, 35 nucleotides, 40 nucleotides, and 50 nucleotides in length. Of course, cross-hybridizing nucleic acids that hybridize to a larger portion of the reference nucleic acid - for example, to a portion of at least 50 nt, at least 100 nt, at least 150 nt, 200 nt, 250 nt, 300 nt, 350 nt, 400 nt, 450
15 nt, or 500 nt or more - or even to the entire length of the reference nucleic acid, are also useful.

 The hybridizing portion of the cross-hybridizing nucleic acid is at least 75% identical in sequence to at least a portion of the reference nucleic acid. Typically, the
20 hybridizing portion of the cross-hybridizing nucleic acid is at least 80%, often at least 85%, 86%, 87%, 88%, 89% or even at least 90% identical in sequence to at least a portion of the reference nucleic acid. Often, the hybridizing portion of the cross-hybridizing nucleic acid will be at least 91%, 92%, 93%,
25 94%, 95%, 96%, 97%, 98%, or 99% identical in sequence to at least a portion of the reference nucleic acid sequence. At times, the hybridizing portion of the cross-hybridizing nucleic acid will be at least 99.5% identical in sequence to at least a portion of the reference nucleic acid.

30 The invention also provides fragments of various of the isolated nucleic acids of the present invention.

 By "fragments" of a reference nucleic acid is here intended isolated nucleic acids, however obtained, that have a

nucleotide sequence identical to a portion of the reference nucleic acid sequence, which portion is at least 17 nucleotides and less than the entirety of the reference nucleic acid. As so defined, "fragments" need not be obtained by physical fragmentation of the reference nucleic acid, although such provenance is not thereby precluded.

In theory, an oligonucleotide of 17 nucleotides is of sufficient length as to occur at random less frequently than once in the three gigabase human genome, and thus to provide a nucleic acid probe that can uniquely identify the reference sequence in a nucleic acid mixture of genomic complexity. As is well known, further specificity can be obtained by probing nucleic acid samples of subgenomic complexity, and/or by using plural fragments as short as 17 nucleotides in length collectively to prime amplification of nucleic acids, as, e.g., by polymerase chain reaction (PCR).

As further described herein below, nucleic acid fragments that encode at least 6 contiguous amino acids (i.e., fragments of 18 nucleotides or more) are useful in directing the expression or the synthesis of peptides that have utility in mapping the epitopes of the protein encoded by the reference nucleic acid. See, e.g., Geysen et al., "Use of peptide synthesis to probe viral antigens for epitopes to a resolution of a single amino acid," Proc. Natl. Acad. Sci. USA 81:3998-4002 (1984); and U.S. Pat. Nos. 4,708,871 and 5,595,915, the disclosures of which are incorporated herein by reference in their entireties.

As further described herein below, fragments that encode at least 8 contiguous amino acids (i.e., fragments of 24 nucleotides or more) are useful in directing the expression or the synthesis of peptides that have utility as immunogens. See, e.g., Lerner, "Tapping the immunological repertoire to produce antibodies of predetermined specificity," Nature

299:592-596 (1982); Shinnick et al., "Synthetic peptide immunogens as vaccines," *Annu. Rev. Microbiol.* 37:425-46 (1983); Sutcliffe et al., "Antibodies that react with predetermined sites on proteins," *Science* 219:660-6 (1983), the disclosures of which are incorporated herein by reference in their entireties.

The nucleic acid fragment of the present invention is thus at least 17 nucleotides in length, typically at least 18 nucleotides in length, and often at least 24 nucleotides in length. Often, the nucleic acid of the present invention is at least 25 nucleotides in length, and even 30 nucleotides, 35 nucleotides, 40 nucleotides, or 45 nucleotides in length. Of course, larger fragments having at least 50 nt, at least 100 nt, at least 150 nt, 200 nt, 250 nt, 300 nt, 350 nt, 400 nt, 450 nt, or 500 nt or more are also useful, and at times preferred.

Having been based upon the mining of genomic sequence, rather than upon surveillance of expressed message, the present invention further provides isolated genome-derived nucleic acids that include portions of the RGL3 gene.

The invention particularly provides genome-derived single exon probes.

As further described in commonly owned and copending U.S. patent application serial nos. 09/864,761, filed May 23, 2001; 09/774,203, filed January 29, 2001; and 09/632,366, filed August 3, 2000, the disclosures of which are incorporated herein by reference in their entireties, "a single exon probe" comprises at least part of an exon ("reference exon") and can hybridize detectably under high stringency conditions to transcript-derived nucleic acids that include the reference exon. The single exon probe will not, however, hybridize detectably under high stringency conditions to nucleic acids that lack the reference exon and instead consist of one or more

exons that are found adjacent to the reference exon in the genome.

Genome-derived single exon probes typically further comprise, contiguous to a first end of the exon portion, a
5 first intronic and/or intergenic sequence that is identically contiguous to the exon in the genome. Often, the genome-derived single exon probe further comprises, contiguous to a second end of the exonic portion, a second intronic and/or intergenic sequence that is identically contiguous to the exon
10 in the genome.

The minimum length of genome-derived single exon probes is defined by the requirement that the exonic portion be of sufficient length to hybridize under high stringency conditions to transcript-derived nucleic acids. Accordingly,
15 the exon portion is at least 17 nucleotides, typically at least 18 nucleotides, 20 nucleotides, 24 nucleotides, 25 nucleotides or even 30, 35, 40, 45, or 50 nucleotides in length, and can usefully include the entirety of the exon, up to 100 nt, 150 nt, 200 nt, 250 nt, 300 nt, 350 nt, 400 nt or even 500 nt or
20 more in length.

The maximum length of genome-derived single exon probes is defined by the requirement that the probes contain portions of no more than one exon, that is, be unable to hybridize detectably under high stringency conditions to
25 nucleic acids that lack the reference exon but include one or more exons that are found adjacent to the reference exon in the genome.

Given variable spacing of exons through eukaryotic genomes, the maximum length of single exon probes of the present invention is typically no more than 25 kb, often no more than 20 kb, 15 kb, 10 kb or 7.5 kb, or even no more than 5 kb, 4 kb, 3 kb, or even no more than about 2.5 kb in length.

The genome-derived single exon probes of the present invention can usefully include at least a first terminal priming sequence not found in contiguity with the rest of the probe sequence in the genome, and often will contain a second 5 terminal priming sequence not found in contiguity with the rest of the probe sequence in the genome.

The present invention also provides isolated genome-derived nucleic acids that include nucleic acid sequence elements that control transcription of the RGL3 gene.

10 With a complete draft of the human genome now available, genomic sequences that are within the vicinity of the RGL3 coding region (and that are additional to those described with particularity herein) can readily be obtained by PCR amplification.

15 The isolated nucleic acids of the present invention can be composed of natural nucleotides in native 5'-3' phosphodiester internucleoside linkage - e.g., DNA or RNA - or can contain any or all of nonnatural nucleotide analogues, nonnative internucleoside bonds, or post-synthesis 20 modifications, either throughout the length of the nucleic acid or localized to one or more portions thereof.

As is well known in the art, when the isolated nucleic acid is used as a hybridization probe, the range of such nonnatural analogues, nonnative internucleoside bonds, or 25 post-synthesis modifications will be limited to those that permit sequence-discriminating basepairing of the resulting nucleic acid. When used to direct expression or RNA or protein *in vitro* or *in vivo*, the range of such nonnatural analogues, nonnative internucleoside bonds, or post-synthesis 30 modifications will be limited to those that permit the nucleic acid to function properly as a polymerization substrate. When the isolated nucleic acid is used as a therapeutic agent, the

range of such changes will be limited to those that do not confer toxicity upon the isolated nucleic acid.

For example, when desired to be used as probes, the isolated nucleic acids of the present invention can usefully 5 include nucleotide analogues that incorporate labels that are directly detectable, such as radiolabels or fluorophores, or nucleotide analogues that incorporate labels that can be visualized in a subsequent reaction, such as biotin or various haptens.

Common radiolabeled analogues include those labeled 10 with ^{33}P , ^{32}P , and ^{35}S , such as α - ^{32}P -dATP, α - ^{32}P -dCTP, α - ^{32}P -dGTP, α - ^{32}P -dTTP, α - ^{32}P -3'-dATP, α - ^{32}P -ATP, α - ^{32}P -CTP, α - ^{32}P -GTP, α - ^{32}P -UTP, α - ^{35}S -dATP, γ - ^{35}S -GTP, γ - ^{33}P -dATP, and the like.

Commercially available fluorescent nucleotide 15 analogues readily incorporated into the nucleic acids of the present invention include Cy3-dCTP, Cy3-dUTP, Cy5-dCTP, Cy3-dUTP (Amersham Pharmacia Biotech, Piscataway, New Jersey, USA), fluorescein-12-dUTP, tetramethylrhodamine-6-dUTP, Texas Red®-5-dUTP, Cascade Blue®-7-dUTP, BODIPY® FL-14-dUTP, BODIPY® 20 TMR-14-dUTP, BODIPY® TR-14-dUTP, Rhodamine Green™-5-dUTP, Oregon Green® 488-5-dUTP, Texas Red®-12-dUTP, BODIPY® 630/650-14-dUTP, BODIPY® 650/665-14-dUTP, Alexa Fluor® 488-5-dUTP, Alexa Fluor® 532-5-dUTP, Alexa Fluor® 568-5-dUTP, Alexa Fluor® 594-5-dUTP, Alexa Fluor® 546-14-dUTP, 25 fluorescein-12-UTP, tetramethylrhodamine-6-UTP, Texas Red®-5-UTP, Cascade Blue®-7-UTP, BODIPY® FL-14-UTP, BODIPY® TMR-14-UTP, BODIPY® TR-14-UTP, Rhodamine Green™-5-UTP, Alexa Fluor® 488-5-UTP, Alexa Fluor® 546-14-UTP (Molecular Probes, Inc. Eugene, OR, USA).

Protocols are available for custom synthesis of 30 nucleotides having other fluorophores. Henegariu *et al.*, "Custom Fluorescent-Nucleotide Synthesis as an Alternative Method for Nucleic Acid Labeling," *Nature Biotechnol.* 18:345 -

348 (2000), the disclosure of which is incorporated herein by reference in its entirety.

Haptens that are commonly conjugated to nucleotides for subsequent labeling include biotin (biotin-11-dUTP, 5 Molecular Probes, Inc., Eugene, OR, USA; biotin-21-UTP, biotin-21-dUTP, Clontech Laboratories, Inc., Palo Alto, CA, USA), digoxigenin (DIG-11-dUTP, alkali labile, DIG-11-UTP, Roche Diagnostics Corp., Indianapolis, IN, USA), and dinitrophenyl (dinitrophenyl-11-dUTP, Molecular Probes, Inc., Eugene, OR, 10 USA).

As another example, when desired to be used for antisense inhibition of transcription or translation, the isolated nucleic acids of the present invention can usefully include altered, often nuclease-resistant, internucleoside bonds. See Hartmann et al. (eds.), Manual of Antisense Methodology (Perspectives in Antisense Science), Kluwer Law International (1999) (ISBN:079238539X); Stein et al. (eds.), Applied Antisense Oligonucleotide Technology, Wiley-Liss (cover 15 (1998) (ISBN: 0471172790); Chadwick et al. (eds.), Oligonucleotides as Therapeutic Agents - Symposium No. 209, John Wiley & Son Ltd (1997) (ISBN: 0471972797), the disclosures 20 of which are incorporated herein by reference in their entireties. Such altered internucleoside bonds are often desired also when the isolated nucleic acid of the present invention is to be used for targeted gene correction, Gamper et al., *Nucl. Acids Res.* 28(21):4332-4339 (2000), the disclosures 25 of which are incorporated herein by reference in its entirety.

Modified oligonucleotide backbones often preferred when the nucleic acid is to be used for antisense purposes are, 30 for example, phosphorothioates, chiral phosphorothioates, phosphorodithioates, phosphotriesters, aminoalkylphosphotriesters, methyl and other alkyl phosphonates including 3'-alkylene phosphonates and chiral phosphonates,

phosphinates, phosphoramidates including 3'-amino phosphoramidate and aminoalkylphosphoramidates, thionophosphoramidates, thionoalkylphosphonates, thionoalkylphosphotriesters, and boranophosphates having normal 5 3'-5' linkages, 2'-5' linked analogs of these, and those having inverted polarity wherein the adjacent pairs of nucleoside units are linked 3'-5' to 5'-3' or 2'-5' to 5'-2'. Representative U.S. patents that teach the preparation of the above phosphorus-containing linkages include, but are not 10 limited to, U.S. Pat. Nos. 3,687,808; 4,469,863; 4,476,301; 5,023,243; 5,177,196; 5,188,897; 5,264,423; 5,276,019; 5,278,302; 5,286,717; 5,321,131; 5,399,676; 5,405,939; 5,453,496; 5,455,233; 5,466,677; 5,476,925; 5,519,126; 5,536,821; 5,541,306; 5,550,111; 5,563,253; 5,571,799; 15 5,587,361; and 5,625,050, the disclosures of which are incorporated herein by reference in their entireties.

Preferred modified oligonucleotide backbones for antisense use that do not include a phosphorus atom have backbones that are formed by short chain alkyl or cycloalkyl 20 internucleoside linkages, mixed heteroatom and alkyl or cycloalkyl internucleoside linkages, or one or more short chain heteroatomic or heterocyclic internucleoside linkages. These include those having morpholino linkages (formed in part from the sugar portion of a nucleoside); siloxane backbones; sulfide, sulfoxide and sulfone backbones; formacetyl and thioformacetyl backbones; methylene formacetyl and thioformacetyl backbones; alkene containing backbones; sulfamate backbones; methyleneimino and methylenehydrazino backbones; sulfonate and sulfonamide backbones; amide 25 backbones; and others having mixed N, O, S and CH₂ component parts. Representative U.S. patents that teach the preparation of the above backbones include, but are not limited to, U.S. Pat. Nos. 5,034,506; 5,166,315; 5,185,444; 5,214,134;

5,216,141; 5,235,033; 5,264,562; 5,264,564; 5,405,938;
5,434,257; 5,466,677; 5,470,967; 5,489,677; 5,541,307;
5,561,225; 5,596,086; 5,602,240; 5,610,289; 5,602,240;
5,608,046; 5,610,289; 5,618,704; 5,623,070; 5,663,312;
5 5,633,360; 5,677,437; and 5,677,439, the disclosures of which
are incorporated herein by reference in their entireties.

In other preferred oligonucleotide mimetics, both the sugar and the internucleoside linkage are replaced with novel groups, such as peptide nucleic acids (PNA).

10 In PNA compounds, the phosphodiester backbone of the nucleic acid is replaced with an amide-containing backbone, in particular by repeating N-(2-aminoethyl) glycine units linked by amide bonds. Nucleobases are bound directly or indirectly to aza nitrogen atoms of the amide portion of the backbone,
15 typically by methylene carbonyl linkages.

The uncharged nature of the PNA backbone provides PNA/DNA and PNA/RNA duplexes with a higher thermal stability than is found in DNA/DNA and DNA/RNA duplexes, resulting from the lack of charge repulsion between the PNA and DNA or RNA
20 strand. In general, the T_m of a PNA/DNA or PNA/RNA duplex is 1°C higher per base pair than the T_m of the corresponding DNA/DNA or DNA/RNA duplex (in 100 mM NaCl).

The neutral backbone also allows PNA to form stable DNA duplexes largely independent of salt concentration. At low
25 ionic strength, PNA can be hybridized to a target sequence at temperatures that make DNA hybridization problematic or impossible. And unlike DNA/DNA duplex formation, PNA hybridization is possible in the absence of magnesium. Adjusting the ionic strength, therefore, is useful if competing
30 DNA or RNA is present in the sample, or if the nucleic acid being probed contains a high level of secondary structure.

PNA also demonstrates greater specificity in binding to complementary DNA. A PNA/DNA mismatch is more destabilizing

than DNA/DNA mismatch. A single mismatch in mixed a PNA/DNA 15-mer lowers the T_m by 8-20°C (15°C on average). In the corresponding DNA/DNA duplexes, a single mismatch lowers the T_m by 4-16°C (11°C on average). Because PNA probes can be 5 significantly shorter than DNA probes, their specificity is greater.

Additionally, nucleases and proteases do not recognize the PNA polyamide backbone with nucleobase sidechains. As a result, PNA oligomers are resistant to 10 degradation by enzymes, and the lifetime of these compounds is extended both *in vivo* and *in vitro*. In addition, PNA is stable over a wide pH range.

Because its backbone is formed from amide bonds, PNA can be synthesized using a modified peptide synthesis protocol. 15 PNA oligomers can be synthesized by both Fmoc and tBoc methods. Representative U.S. patents that teach the preparation of PNA compounds include, but are not limited to, U.S. Pat. Nos. 5,539,082; 5,714,331; and 5,719,262, each of which is herein incorporated by reference; automated PNA 20 synthesis is readily achievable on commercial synthesizers (see, e.g., "PNA User's Guide," Rev. 2, February 1998, Perseptive Biosystems Part No. 60138, Applied Biosystems, Inc., Foster City, CA).

PNA chemistry and applications are reviewed, *inter alia*, in Ray et al., *FASEB J.* 14(9):1041-60 (2000); Nielsen et al., *Pharmacol Toxicol.* 86(1):3-7 (2000); Larsen et al., *Biochim Biophys Acta.* 1489(1):159-66 (1999); Nielsen, *Curr. Opin. Struct. Biol.* 9(3):353-7 (1999), and Nielsen, *Curr. Opin. Biotechnol.* 10(1):71-5 (1999), the disclosures of which are 30 incorporated herein by reference in their entireties.

Differences from nucleic acid compositions found in nature – e.g., nonnative bases, altered internucleoside linkages, post-synthesis modification – can be present

throughout the length of the nucleic acid or can, instead, usefully be localized to discrete portions thereof. As an example of the latter, chimeric nucleic acids can be synthesized that have discrete DNA and RNA domains and 5 demonstrated utility for targeted gene repair, as further described in U.S. Pat. Nos. 5,760,012 and 5,731,181, the disclosures of which are incorporated herein by reference in their entireties. As another example, chimeric nucleic acids comprising both DNA and PNA have been demonstrated to have 10 utility in modified PCR reactions. See Misra et al., *Biochem.* 37: 1917-1925 (1998); see also Finn et al., *Nucl. Acids Res.* 24: 3357-3363 (1996), incorporated herein by reference.

Unless otherwise specified, nucleic acids of the present invention can include any topological conformation 15 appropriate to the desired use; the term thus explicitly comprehends, among others, single-stranded, double-stranded, triplexed, quadruplexed, partially double-stranded, partially-triplexed, partially-quadruplexed, branched, hairpinned, circular, and padlocked conformations. Padlock conformations 20 and their utilities are further described in Banér et al., *Curr. Opin. Biotechnol.* 12:11-15 (2001); Escude et al., *Proc. Natl. Acad. Sci. USA* 14;96(19):10603-7 (1999); Nilsson et al., *Science* 265(5181):2085-8 (1994), the disclosures of which are incorporated herein by reference in their entireties. Triplex 25 and quadruplex conformations, and their utilities, are reviewed in Praseuth et al., *Biochim. Biophys. Acta* 1489(1):181-206 (1999); Fox, *Curr. Med. Chem.* 7(1):17-37 (2000); Kochetkova et al., *Methods Mol. Biol.* 130:189-201 (2000); Chan et al., *J. Mol. Med.* 75(4):267-82 (1997), the disclosures of which are 30 incorporated herein by reference in their entireties.

The nucleic acids of the present invention can be detectably labeled.

Commonly-used labels include radionuclides, such as ^{32}P , ^{33}P , ^{35}S , ^3H (and for NMR detection, ^{13}C and ^{15}N), haptens that can be detected by specific antibody or high affinity binding partner (such as avidin), and fluorophores.

5 As noted above, detectable labels can be incorporated by inclusion of labeled nucleotide analogues in the nucleic acid. Such analogues can be incorporated by enzymatic polymerization, such as by nick translation, random priming, polymerase chain reaction (PCR), terminal transferase tailing, 10 and end-filling of overhangs, for DNA molecules, and *in vitro* transcription driven, e.g., from phage promoters, such as T7, T3, and SP6, for RNA molecules. Commercial kits are readily available for each such labeling approach.

Analogues can also be incorporated during automated 15 solid phase chemical synthesis.

As is well known, labels can also be incorporated after nucleic acid synthesis, with the 5' phosphate and 3' hydroxyl providing convenient sites for post-synthetic covalent attachment of detectable labels.

20 Various other post-synthetic approaches permit internal labeling of nucleic acids.

For example, fluorophores can be attached using a cisplatin reagent that reacts with the N7 of guanine residues (and, to a lesser extent, adenine bases) in DNA, RNA, and PNA 25 to provide a stable coordination complex between the nucleic acid and fluorophore label (Universal Linkage System) (available from Molecular Probes, Inc., Eugene, OR, USA and Amersham Pharmacia Biotech, Piscataway, NJ, USA); see Alers et al., *Genes, Chromosomes & Cancer*, Vol. 25, pp. 301 - 305 30 (1999); Jelsma et al., *J. NIH Res.* 5:82 (1994); Van Belkum et al., *BioTechniques* 16:148-153 (1994), incorporated herein by reference. As another example, nucleic acids can be labeled using a disulfide-containing linker (FastTagTM Reagent, Vector

Laboratories, Inc., Burlingame, CA, USA) that is photo- or thermally coupled to the target nucleic acid using aryl azide chemistry; after reduction, a free thiol is available for coupling to a hapten, fluorophore, sugar, affinity ligand, or 5 other marker.

Multiple independent or interacting labels can be incorporated into the nucleic acids of the present invention.

For example, both a fluorophore and a moiety that in proximity thereto acts to quench fluorescence can be included 10 to report specific hybridization through release of fluorescence quenching, Tyagi et al., *Nature Biotechnol.* 14: 303-308 (1996); Tyagi et al., *Nature Biotechnol.* 16, 49-53 (1998); Sokol et al., *Proc. Natl. Acad. Sci. USA* 95: 11538-11543 (1998); Kostrikis et al., *Science* 279:1228-1229 (1998); Marras et al., *Genet. Anal.* 14: 151-156 (1999); U.S. 15 Pat. Nos. 5,846,726, 5,925,517, 5925517, or to report exonucleotidic excision, U.S. Pat. No. 5,538,848; Holland et al., *Proc. Natl. Acad. Sci. USA* 88:7276-7280 (1991); Heid et al., *Genome Res.* 6(10):986-94 (1996); Kuimelis et al., *Nucleic 20 Acids Symp Ser.* (37):255-6 (1997); U.S. Patent No. 5,723,591, the disclosures of which are incorporated herein by reference in their entireties.

So labeled, the isolated nucleic acids of the present invention can be used as probes, as further described below.

25 Nucleic acids of the present invention can also usefully be bound to a substrate. The substrate can porous or solid, planar or non-planar, unitary or distributed; the bond can be covalent or noncovalent. Bound to a substrate, nucleic acids of the present invention can be used as probes in their 30 unlabeled state.

For example, the nucleic acids of the present invention can usefully be bound to a porous substrate, commonly a membrane, typically comprising nitrocellulose, nylon, or

positively-charged derivatized nylon; so attached, the nucleic acids of the present invention can be used to detect RGL3 nucleic acids present within a labeled nucleic acid sample, either a sample of genomic nucleic acids or a sample of 5 transcript-derived nucleic acids, e.g. by reverse dot blot.

The nucleic acids of the present invention can also usefully be bound to a solid substrate, such as glass, although other solid materials, such as amorphous silicon, crystalline silicon, or plastics, can also be used. Such plastics include 10 polymethylacrylic, polyethylene, polypropylene, polyacrylate, polymethylmethacrylate, polyvinylchloride, polytetrafluoroethylene, polystyrene, polycarbonate, polyacetal, polysulfone, celluloseacetate, cellulosenitrate, nitrocellulose, or mixtures thereof.

15 Typically, the solid substrate will be rectangular, although other shapes, particularly disks and even spheres, present certain advantages. Particularly advantageous alternatives to glass slides as support substrates for array of nucleic acids are optical discs, as described in Demers, 20 "Spatially Addressable Combinatorial Chemical Arrays in CD-ROM Format," international patent publication WO 98/12559, incorporated herein by reference in its entirety.

The nucleic acids of the present invention can be attached covalently to a surface of the support substrate or 25 applied to a derivatized surface in a chaotropic agent that facilitates denaturation and adherence by presumed noncovalent interactions, or some combination thereof.

The nucleic acids of the present invention can be bound to a substrate to which a plurality of other nucleic acids are concurrently bound, hybridization to each of the 30 plurality of bound nucleic acids being separately detectable. At low density, e.g. on a porous membrane, these substrate-bound collections are typically denominated macroarrays; at

higher density, typically on a solid support, such as glass, these substrate bound collections of plural nucleic acids are colloquially termed microarrays. As used herein, the term microarray includes arrays of all densities. It is, therefore, 5 another aspect of the invention to provide microarrays that include the nucleic acids of the present invention.

The isolated nucleic acids of the present invention can be used as hybridization probes to detect, characterize, and quantify RGL3 nucleic acids in, and isolate RGL3 nucleic 10 acids from, both genomic and transcript-derived nucleic acid samples. When free in solution, such probes are typically, but not invariably, detectably labeled; bound to a substrate, as in a microarray, such probes are typically, but not invariably unlabeled.

15 For example, the isolated nucleic acids of the present invention can be used as probes to detect and characterize gross alterations in the RGL3 genomic locus, such as deletions, insertions, translocations, and duplications of the RGL3 genomic locus through fluorescence *in situ* hybridization (FISH) to chromosome spreads. See, e.g.,
20 Andreeff et al. (eds.), Introduction to Fluorescence In Situ Hybridization: Principles and Clinical Applications, John Wiley & Sons (1999) (ISBN: 0471013455), the disclosure of which is incorporated herein by reference in its entirety. The isolated
25 nucleic acids of the present invention can be used as probes to assess smaller genomic alterations using, e.g., Southern blot detection of restriction fragment length polymorphisms. The isolated nucleic acids of the present invention can be used as probes to isolate genomic clones that include the nucleic acids
30 of the present invention, which thereafter can be restriction mapped and sequenced to identify deletions, insertions, translocations, and substitutions (single nucleotide polymorphisms, SNPs) at the sequence level.

The isolated nucleic acids of the present invention can also be used as probes to detect, characterize, and quantify RGL3 nucleic acids in, and isolate RGL3 nucleic acids from, transcript-derived nucleic acid samples.

For example, the isolated nucleic acids of the present invention can be used as hybridization probes to detect, characterize by length, and quantify RGL3 mRNA by northern blot of total or poly-A⁺- selected RNA samples. For example, the isolated nucleic acids of the present invention can be used as hybridization probes to detect, characterize by location, and quantify RGL3 message by *in situ* hybridization to tissue sections (see, e.g., Schwarchzacher et al., In Situ Hybridization, Springer-Verlag New York (2000) (ISBN: 0387915966), the disclosure of which is incorporated herein by reference in its entirety). For example, the isolated nucleic acids of the present invention can be used as hybridization probes to measure the representation of RGL3 clones in a cDNA library. For example, the isolated nucleic acids of the present invention can be used as hybridization probes to isolate RGL3 nucleic acids from cDNA libraries, permitting sequence level characterization of RGL3 messages, including identification of deletions, insertions, truncations – including deletions, insertions, and truncations of exons in alternatively spliced forms – and single nucleotide polymorphisms.

All of the aforementioned probe techniques are well within the skill in the art, and are described at greater length in standard texts such as Sambrook et al., Molecular Cloning: A Laboratory Manual (3rd ed.), Cold Spring Harbor Laboratory Press (2001) (ISBN: 0879695773); Ausubel et al. (eds.), Short Protocols in Molecular Biology: A Compendium of Methods from Current Protocols in Molecular Biology (4th ed.), John Wiley & Sons, 1999 (ISBN: 047132938X); and Walker et al.

(eds.), The Nucleic Acids Protocols Handbook, Humana Press (2000) (ISBN: 0896034593), the disclosures of which are incorporated herein by reference in their entirety.

As described in the Examples herein below, the nucleic acids of the present invention can also be used to detect and quantify RGL3 nucleic acids in transcript-derived samples – that is, to measure expression of the RGL3 gene – when included in a microarray. Measurement of RGL3 expression has particular utility in the diagnosis and treatment of cancer, as further described in the Examples herein below.

As would be readily apparent to one of skill in the art, each RGL3 nucleic acid probe – whether labeled, substrate-bound, or both – is thus currently available for use as a tool for measuring the level of RGL3 expression in each of the tissues in which expression has already been confirmed, notably adrenal, adult liver, bone marrow, brain, fetal liver, heart, kidney, lung, placenta, colon, skeletal muscle and prostate. The utility is specific to the probe: under high stringency conditions, the probe reports the level of expression of message specifically containing that portion of the RGL3 gene included within the probe.

Measuring tools are well known in many arts, not just in molecular biology, and are known to possess credible, specific, and substantial utility. For example, U.S. Patent No. 6,016,191 describes and claims a tool for measuring characteristics of fluid flow in a hydrocarbon well; U.S. Patent No. 6,042,549 describes and claims a device for measuring exercise intensity; U.S. Patent No. 5,889,351 describes and claims a device for measuring viscosity and for measuring characteristics of a fluid; U.S. Patent No. 5,570,694 describes and claims a device for measuring blood pressure; U.S. Patent No. 5,930,143 describes and claims a device for measuring the dimensions of machine tools; U.S. Patent No.

5,279,044 describes and claims a measuring device for determining an absolute position of a movable element; U.S. Patent No. 5,186,042 describes and claims a device for measuring action force of a wheel; and U.S. Patent No. 5 4,246,774 describes and claims a device for measuring the draft of smoking articles such as cigarettes.

As for tissues not yet demonstrated to express RGL3, the RGL3 nucleic acid probes of the present invention are currently available as tools for surveying such tissues to 10 detect the presence of RGL3 nucleic acids.

Survey tools - i.e., tools for determining the presence and/or location of a desired object by search of an area - are well known in many arts, not just in molecular biology, and are known to possess credible, specific, and 15 substantial utility. For example, U.S. Patent No. 6,046,800 describes and claims a device for surveying an area for objects that move; U.S. Patent No. 6,025,201 describes and claims an apparatus for locating and discriminating platelets from non-platelet particles or cells on a cell-by-cell basis in a 20 whole blood sample; U.S. Patent No. 5,990,689 describes and claims a device for detecting and locating anomalies in the electromagnetic protection of a system; U.S. Patent No. 5,984,175 describes and claims a device for detecting and identifying wearable user identification units; U.S. Patent No. 25 3,980,986 ("Oil well survey tool"), describes and claims a tool for finding the position of a drill bit working at the bottom of a borehole.

As noted above, the nucleic acid probes of the present invention are useful in constructing microarrays; the 30 microarrays, in turn, are products of manufacture that are useful for measuring and for surveying gene expression.

When included on a microarray, each RGL3 nucleic acid probe makes the microarray specifically useful for detecting

that portion of the RGL3 gene included within the probe, thus imparting upon the microarray device the ability to detect a signal where, absent such probe, it would have reported no signal. This utility makes each individual probe on such 5 microarray akin to an antenna, circuit, firmware or software element included in an electronic apparatus, where the antenna, circuit, firmware or software element imparts upon the apparatus the ability newly and additionally to detect signal in a portion of the radio-frequency spectrum where previously 10 it could not; such devices are known to have specific, substantial, and credible utility.

Changes in the level of expression need not be observed for the measurement of expression to have utility.

For example, where gene expression analysis is used 15 to assess toxicity of chemical agents on cells, the failure of the agent to change a gene's expression level is evidence that the drug likely does not affect the pathway of which the gene's expressed protein is a part. Analogously, where gene expression analysis is used to assess side effects of 20 pharmacologic agents – whether in lead compound discovery or in subsequent screening of lead compound derivatives – the inability of the agent to alter a gene's expression level is evidence that the drug does not affect the pathway of which the gene's expressed protein is a part.

WO 99/58720, incorporated herein by reference in its 25 entirety, provides methods for quantifying the relatedness of a first and second gene expression profile and for ordering the relatedness of a plurality of gene expression profiles, without regard to the identity or function of the genes whose 30 expression is used in the calculation.

Gene expression analysis, including gene expression analysis by microarray hybridization, is, of course, principally a laboratory-based art. Devices and apparatus used

principally in laboratories to facilitate laboratory research
are well-established to possess specific, substantial, and
credible utility. For example, U.S. Patent No. 6,001,233
describes and claims a gel electrophoresis apparatus having a
5 cam-activated clamp; for example, U.S. Patent No. 6,051,831
describes and claims a high mass detector for use in time-of-
flight mass spectrometers; for example, U.S. Patent NO.
5,824,269 describes and claims a flow cytometer-as is well
known, few gel electrophoresis apparatuses, TOF-MS devices, or
10 flow cytometers are sold for consumer use.

Indeed, and in particular, nucleic acid microarrays,
as devices intended for laboratory use in measuring gene
expression, are well-established to have specific, substantial
and credible utility. Thus, the microarrays of the present
15 invention have at least the specific, substantial and credible
utilities of the microarrays claimed as devices and articles of
manufacture in the following U.S. patents, the disclosures of
each of which is incorporated herein by reference: U.S. Patent
Nos. 5,445,934 ("Array of oligonucleotides on a solid
20 substrate"); 5,744,305 ("Arrays of materials attached to a
substrate"); and 6,004,752 ("Solid support with attached
molecules").

Genome-derived single exon probes and genome-derived
single exon probe microarrays have the additional utility,
25 *inter alia*, of permitting high-throughput detection of splice
variants of the nucleic acids of the present invention, as
further described in copending and commonly owned U.S. Patent
application no. 09/632,366, filed August 3, 2000, the
disclosure of which is incorporated herein by reference in its
30 entirety.

The isolated nucleic acids of the present invention
can also be used to prime synthesis of nucleic acid, for

purpose of either analysis or isolation, using mRNA, cDNA, or genomic DNA as template.

For use as primers, at least 17 contiguous nucleotides of the isolated nucleic acids of the present invention will be used. Often, at least 18, 19, or 20 contiguous nucleotides of the nucleic acids of the present invention will be used, and on occasion at least 20, 22, 24, or 25 contiguous nucleotides of the nucleic acids of the present invention will be used, and even 30 nucleotides or more of the nucleic acids of the present invention can be used to prime specific synthesis.

The nucleic acid primers of the present invention can be used, for example, to prime first strand cDNA synthesis on an mRNA template.

Such primer extension can be done directly to analyze the message. Alternatively, synthesis on an mRNA template can be done to produce first strand cDNA. The first strand cDNA can thereafter be used, *inter alia*, directly as a single-stranded probe, as above-described, as a template for sequencing — permitting identification of alterations, including deletions, insertions, and substitutions, both normal allelic variants and mutations associated with abnormal phenotypes— or as a template, either for second strand cDNA synthesis (e.g., as an antecedent to insertion into a cloning or expression vector), or for amplification.

The nucleic acid primers of the present invention can also be used, for example, to prime single base extension (SBE) for SNP detection (see, e.g., U.S. Pat. No. 6,004,744, the disclosure of which is incorporated herein by reference in its entirety).

As another example, the nucleic acid primers of the present invention can be used to prime amplification of RGL3

nucleic acids, using transcript-derived or genomic DNA as template.

Primer-directed amplification methods are now well-established in the art. Methods for performing the polymerase chain reaction (PCR) are compiled, *inter alia*, in McPherson, PCR (Basics: From Background to Bench), Springer Verlag (2000) (ISBN: 0387916008); Innis et al. (eds.), PCR Applications: Protocols for Functional Genomics, Academic Press (1999) (ISBN: 0123721857); Gelfand et al. (eds.), PCR Strategies, Academic Press (1998) (ISBN: 0123721822); Newton et al., PCR, Springer-Verlag New York (1997) (ISBN: 0387915060); Burke (ed.), PCR: Essential Techniques, John Wiley & Son Ltd (1996) (ISBN: 047195697X); White (ed.), PCR Cloning Protocols: From Molecular Cloning to Genetic Engineering, Vol. 67, Humana Press (1996) (ISBN: 0896033430); McPherson et al. (eds.), PCR 2: A Practical Approach, Oxford University Press, Inc. (1995) (ISBN: 0199634254), the disclosures of which are incorporated herein by reference in their entireties. Methods for performing RT-PCR are collected, e.g., in Siebert et al. (eds.), Gene Cloning and Analysis by RT-PCR, Eaton Publishing Company/Bio Techniques Books Division, 1998 (ISBN: 1881299147); Siebert (ed.), PCR Technique:RT-PCR, Eaton Publishing Company/BioTechniques Books (1995) (ISBN:1881299139), the disclosure of which is incorporated herein by reference in its entirety.

Isothermal amplification approaches, such as rolling circle amplification, are also now well-described. See, e.g., Schweitzer et al., *Curr. Opin. Biotechnol.* 12(1):21-7 (2001); U.S. Patent Nos. 6,235,502, 6,221,603, 6,210,884, 6,183,960, 5,854,033, 5,714,320, 5,648,245, and international patent publications WO 97/19193 and WO 00/15779, the disclosures of which are incorporated herein by reference in their entireties. Rolling circle amplification can be combined with other

techniques to facilitate SNP detection. See, e.g., Lizardi et al., *Nature Genet.* 19(3):225-32 (1998).

As further described below, nucleic acids of the present invention, inserted into vectors that flank the nucleic acid insert with a phage promoter, such as T7, T3, or SP6 promoter, can be used to drive *in vitro* expression of RNA complementary to either strand of the nucleic acid of the present invention. The RNA can be used, *inter alia*, as a single-stranded probe, in cDNA-mRNA subtraction, or for *in vitro* translation.

As will be further discussed herein below, nucleic acids of the present invention that encode RGL3 protein or portions thereof can be used, *inter alia*, to express the RGL3 proteins or protein fragments, either alone, or as part of fusion proteins.

Expression can be from genomic nucleic acids of the present invention, or from transcript-derived nucleic acids of the present invention.

Where protein expression is effected from genomic DNA, expression will typically be effected in eukaryotic, typically mammalian, cells capable of splicing introns from the initial RNA transcript. Expression can be driven from episomal vectors, such as EBV-based vectors, or can be effected from genomic DNA integrated into a host cell chromosome. As will be more fully described below, where expression is from transcript-derived (or otherwise intron-less) nucleic acids of the present invention, expression can be effected in wide variety of prokaryotic or eukaryotic cells.

Expressed *in vitro*, the protein, protein fragment, or protein fusion can thereafter be isolated, to be used, *inter alia*, as a standard in immunoassays specific for the proteins, or protein isoforms, of the present invention; to be used as a therapeutic agent, e.g., to be administered as passive

replacement therapy in individuals deficient in the proteins of the present invention, or to be administered as a vaccine; to be used for *in vitro* production of specific antibody, the antibody thereafter to be used, e.g., as an analytical reagent 5 for detection and quantitation of the proteins of the present invention or to be used as an immunotherapeutic agent.

The isolated nucleic acids of the present invention can also be used to drive *in vivo* expression of the proteins of the present invention. *In vivo* expression can be driven from a 10 vector - typically a viral vector, often a vector based upon a replication incompetent retrovirus, an adenovirus, or an adeno-associated virus (AAV) - for purpose of gene therapy. *In vivo* expression can also be driven from signals endogenous to the nucleic acid or from a vector, often a plasmid vector, such as 15 pVAX1 (Invitrogen, Carlsbad CA, USA), for purpose of "naked" nucleic acid vaccination, as further described in U.S. Pat. Nos. 5,589,466; 5,679,647; 5,804,566; 5,830,877; 5,843,913; 5,880,104; 5,958,891; 5,985,847; 6,017,897; 6,110,898; 6,204,250, the disclosures of which are incorporated herein by 20 reference in their entireties.

The nucleic acids of the present invention can also be used for antisense inhibition of transcription or translation. See Phillips (ed.), Antisense Technology, Part B, Methods in Enzymology Vol. 314, Academic Press, Inc. (1999) 25 (ISBN: 012182215X); Phillips (ed.), Antisense Technology, Part A, Methods in Enzymology Vol. 313, Academic Press, Inc. (1999) (ISBN: 0121822141); Hartmann et al. (eds.), Manual of Antisense Methodology (Perspectives in Antisense Science), Kluwer Law International (1999) (ISBN:079238539X); Stein et al. (eds.), 30 Applied Antisense Oligonucleotide Technology, Wiley-Liss (cover (1998) (ISBN: 0471172790); Agrawal et al. (eds.), Antisense Research and Application, Springer-Verlag New York, Inc. (1998) (ISBN: 3540638334); Lichtenstein et al. (eds.), Antisense

Technology: A Practical Approach, Vol. 185, Oxford University Press, INC. (1998) (ISBN: 0199635838); Gibson (ed.), Antisense and Ribozyme Methodology: Laboratory Companion, Chapman & Hall (1997) (ISBN: 3826100794); Chadwick et al. (eds.),
5 Oligonucleotides as Therapeutic Agents - Symposium No. 209, John Wiley & Son Ltd (1997) (ISBN: 0471972797), the disclosures of which are incorporated herein by reference in their entireties.

Nucleic acids of the present invention, particularly
10 cDNAs of the present invention, that encode full-length human RGL3 protein isoforms, have additional, well-recognized, immediate, real world utility as commercial products of manufacture suitable for sale.

For example, Invitrogen Corp. (Carlsbad, CA, USA),
15 through its Research Genetics subsidiary, sells full length human cDNAs cloned into one of a selection of expression vectors as GeneStorm® expression-ready clones; utility is specific for the gene, since each gene is capable of being ordered separately and has a distinct catalogue number, and
20 utility is substantial, each clone selling for \$650.00 US. Similarly, Incyte Genomics (Palo Alto, CA, USA) sells clones from public and proprietary sources in multi-well plates or individual tubes.

Nucleic acids of the present invention that include
25 genomic regions encoding the human RGL3 protein, or portions thereof, have yet further utilities.

For example, genomic nucleic acids of the present invention can be used as amplification substrates, e.g. for preparation of genome-derived single exon probes of the present
30 invention, as described above and in copending and commonly-owned U.S. patent application nos. 09/864,761, filed May 23, 2001, 09/774,203, filed January 29, 2001, and 09/632,366, filed

August 3, 2000, the disclosures of which are incorporated herein by reference in their entireties.

As another example, genomic nucleic acids of the present invention can be integrated non-homologously into the genome of somatic cells, e.g. CHO cells, COS cells, or 293 cells, with or without amplification of the insertional locus, in order, e.g., to create stable cell lines capable of producing the proteins of the present invention.

As another example, more fully described herein below, genomic nucleic acids of the present invention can be integrated nonhomologously into embryonic stem (ES) cells to create transgenic non-human animals capable of producing the proteins of the present invention.

Genomic nucleic acids of the present invention can also be used to target homologous recombination to the human RGL3 locus. See, e.g., U.S. Patent Nos. 6,187,305; 6,204,061; 5,631,153; 5,627,059; 5,487,992; 5,464,764; 5,614,396; 5,527,695 and 6,063,630; and Kmiec et al. (eds.), Gene Targeting Protocols, Vol. 133, Humana Press (2000) (ISBN: 0896033600); Joyner (ed.), Gene Targeting: A Practical Approach, Oxford University Press, Inc. (2000) (ISBN: 0199637938); Sedivy et al., Gene Targeting, Oxford University Press (1998) (ISBN: 071677013X); Tymms et al. (eds.), Gene Knockout Protocols, Humana Press (2000) (ISBN: 0896035727); Mak et al. (eds.), The Gene Knockout FactsBook, Vol. 2, Academic Press, Inc. (1998) (ISBN: 0124660444); Torres et al., Laboratory Protocols for Conditional Gene Targeting, Oxford University Press (1997) (ISBN: 019963677X); Vega (ed.), Gene Targeting, CRC Press, LLC (1994) (ISBN: 084938950X), the disclosures of which are incorporated herein by reference in their entireties.

Where the genomic region includes transcription regulatory elements, homologous recombination can be used to

alter the expression of RGL3, both for purpose of *in vitro* production of RGL3 protein from human cells, and for purpose of gene therapy. See, e.g., U.S. Pat. Nos. 5,981,214, 6,048,524; 5,272,071.

5 Fragments of the nucleic acids of the present invention smaller than those typically used for homologous recombination can also be used for targeted gene correction or alteration, possibly by cellular mechanisms different from those engaged during homologous recombination.

10 For example, partially duplexed RNA/DNA chimeras have been shown to have utility in targeted gene correction, U.S. Pat. Nos. 5,945,339, 5,888,983, 5,871,984, 5,795,972, 5,780,296, 5,760,012, 5,756,325, 5,731,181, the disclosures of which are incorporated herein by reference in their entireties.

15 So too have small oligonucleotides fused to triplexing domains have been shown to have utility in targeted gene correction, Culver et al., "Correction of chromosomal point mutations in human cells with bifunctional oligonucleotides," *Nature Biotechnol.* 17(10):989-93 (1999), as have oligonucleotides 20 having modified terminal bases or modified terminal internucleoside bonds, Gamper et al., *Nucl. Acids Res.* 28(21):4332-9 (2000), the disclosures of which are incorporated herein by reference.

25 The isolated nucleic acids of the present invention can also be used to provide the initial substrate for recombinant engineering of RGL3 protein variants having desired phenotypic improvements. Such engineering includes, for example, site-directed mutagenesis, random mutagenesis with subsequent functional screening, and more elegant schemes for 30 recombinant evolution of proteins, as are described, *inter alia*, in U.S. Pat. Nos. 6,180,406; 6,165,793; 6,117,679; and 6,096,548, the disclosures of which are incorporated herein by reference in their entireties.

Nucleic acids of the present invention can be obtained by using the labeled probes of the present invention to probe nucleic acid samples, such as genomic libraries, cDNA libraries, and mRNA samples, by standard techniques. Nucleic acids of the present invention can also be obtained by amplification, using the nucleic acid primers of the present invention, as further demonstrated in Example 1, herein below.

Nucleic acids of the present invention of fewer than about 100 nt can also be synthesized chemically, typically by solid phase synthesis using commercially available automated synthesizers.

"Full Length" RGL3 Nucleic Acids

In a first series of nucleic acid embodiments, the invention provides isolated nucleic acids that encode the entirety of the RGL3 protein. As discussed above, the "full-length" nucleic acids of the present invention can be used, *inter alia*, to express full length RGL3 protein. The full-length nucleic acids can also be used as nucleic acid probes; used as probes, the isolated nucleic acids of these embodiments will hybridize to RGL3.

In a first such embodiment, the invention provides an isolated nucleic acid comprising (i) the nucleotide sequence of SEQ ID NO: 1, or (ii) the complement of (i). SEQ ID NO: 1 presents the entire cDNA of RGL3, including the 5' untranslated (UT) region and 3' UT.

In a second embodiment, the invention provides an isolated nucleic acid comprising (i) the nucleotide sequence of SEQ ID NO: 2, (ii) a degenerate variant of the nucleotide sequence of SEQ ID NO: 2, or (iii) the complement of (i) or (ii). SEQ ID NO: 2 presents the open reading frame (ORF) from SEQ ID NO: 1.

In a third embodiment, the invention provides an isolated nucleic acid comprising (i) a nucleotide sequence that encodes a polypeptide with the amino acid sequence of SEQ ID NO: 3 or (ii) the complement of a nucleotide sequence that 5 encodes a polypeptide with the amino acid sequence of SEQ ID NO: 3. SEQ ID NO: 3 provides the amino acid sequence of RGL3.

In a fourth embodiment, the invention provides an isolated nucleic acid having a nucleotide sequence that (i) encodes a polypeptide having the sequence of SEQ ID NO: 3, (ii) 10 encodes a polypeptide having the sequence of SEQ ID NO: 3 with conservative amino acid substitutions, or (iii) that is the complement of (i) or (ii), where SEQ ID NO: 3 provides the amino acid sequence of RGL3.

15

Selected Partial Nucleic Acids

In a second series of nucleic acid embodiments, the invention provides isolated nucleic acids that encode select portions of RGL3. As will be further discussed herein below, 20 these "partial" nucleic acids can be used, *inter alia*, to express specific portions of the RGL3. These "partial" nucleic acids can also be used, *inter alia*, as nucleic probes.

In a first such embodiment, the invention provides an isolated nucleic acid comprising (i) the nucleotide sequence of 25 SEQ ID NO: 4, (ii) a degenerate variant of SEQ ID NO: 4, or (iii) the complement of (i) or (ii), wherein the isolated nucleic acid is no more than about 100 kb in length, typically no more than about 75 kb in length, more typically no more than about 50 kb length. SEQ ID NO: 4 encodes the portion of RGL3 30 which contains a two base pair insertion compared to Genbank entry BC014426. Often, the isolated nucleic acids of this embodiment are no more than about 25 kb in length, often no

more than about 15 kb in length, and frequently no more than about 10 kb in length.

In another embodiment, the invention provides an isolated nucleic acid comprising (i) a nucleotide sequence that encodes SEQ ID NO: 5 or (ii) the complement of a nucleotide sequence that encodes SEQ ID NO: 5, wherein the isolated nucleic acid is no more than about 100 kb in length, typically no more than about 75 kb in length, frequently no more than about 50 kb in length. SEQ ID NO: 5 the amino acid sequence encoded by SEQ ID NO: 4. Often, the isolated nucleic acids of this embodiment are no more than about 25 kb in length, often no more than about 15 kb in length, and frequently no more than about 10 kb in length.

In another embodiment, the invention provides an isolated nucleic acid comprising (i) a nucleotide sequence that encodes SEQ ID NO: 5, (ii) a nucleotide sequence that encodes SEQ ID NO: 5 with conservative substitutions, or (iii) the complement of (i) or (ii), wherein the isolated nucleic acid is no more than about 100 kb in length, typically no more than about 75 kb in length, and often no more than about 50 kb in length. Often, the isolated nucleic acids of this embodiment are no more than about 25 kb in length, often no more than about 15 kb in length, and frequently no more than about 10 kb in length.

25

Cross-Hybridizing Nucleic Acids

In another series of nucleic acid embodiments, the invention provides isolated nucleic acids that hybridize to various of the RGL3 nucleic acids of the present invention. These cross-hybridizing nucleic acids can be used, *inter alia*, as probes for, and to drive expression of, proteins that are

related to RGL3 of the present invention as further isoforms, homologues, paralogues, or orthologues.

In a first embodiment, the invention provides an isolated nucleic acid comprising a sequence that hybridizes under high stringency conditions to a probe the nucleotide sequence of which consists of at least 17 nt, 18, 19, 20, 21, 22, 23, 24, 25, 30, 40, or 50 nt of SEQ ID NO: 4 or the complement of SEQ ID NO: 4, wherein the isolated nucleic acid is no more than about 100 kb in length, typically no more than about 75 kb in length, and often no more than about 50 kb in length. Often, the isolated nucleic acids of this embodiment are no more than about 25 kb in length, often no more than about 15 kb in length, and frequently no more than about 10 kb in length.

In a further embodiment, the invention provides an isolated nucleic acid comprising a sequence that hybridizes under moderate stringency conditions to a probe the nucleotide sequence of which consists of atleast 17 nt, 18, 19, 20, 21, 22, 23, 24, 25, 30, 40, or 50 nt of SEQ ID NO: 4 or the complement of SEQ ID NO: 4, wherein the isolated nucleic acid is no more than about 100 kb in length, typically no more than about 75 kb in length, and often no more than about 50 kb in length. Often, the isolated nucleic acids of this embodiment are no more than about 25 kb in length, often no more than about 15 kb in length, and frequently no more than about 10 kb in length.

In a further embodiment, the invention provides an isolated nucleic acid comprising a sequence that hybridizes under high stringency conditions to a hybridization probe the nucleotide sequence of which (i) encodes a polypeptide having the sequence of SEQ ID NO: 5, (ii) encodes a polypeptide having the sequence of SEQ ID NO: 5 with conservative amino acid substitutions, or (iii) is the complement of (i) or (ii),

wherein the isolated nucleic acid is no more than about 100 kb in length, typically no more than about 75 kb in length, and often no more than about 50 kb in length. Often, the isolated nucleic acids of this embodiment are no more than about 25 kb
5 in length, often no more than about 15 kb in length, and frequently no more than about 10 kb in length.

Particularly Useful Nucleic Acids

10 Particularly useful among the above-described nucleic acids are those that are expressed, or the complement of which are expressed, in adrenal, adult liver, bone marrow, brain, fetal liver, heart, kidney, lung, placenta, colon, skeletal muscle and prostate.

15 Also particularly useful among the above-described nucleic acids are those that encode, or the complement of which encode a guanine nucleotide exchange factor for the small GTPase Ral and a downstream effector for both Rit and Ras.

20 Other particularly useful embodiments of the nucleic acids above-described are those that encode, or the complement of which encode, a polypeptide having any or all of a RasGEFN domain, a RasGEF domain or a RA domain.

Nucleic Acid Fragments

25 In another series of nucleic acid embodiments, the invention provides fragments of various of the isolated nucleic acids of the present invention which prove useful, *inter alia*, as nucleic acid probes, as amplification primers, and to direct expression or synthesis of epitopic or immunogenic protein fragments.

In a further embodiment, the invention provides an isolated nucleic acid comprising at least 17 nucleotides, 18 nucleotides, 20 nucleotides, 24 nucleotides, or 25 nucleotides

of (i) SEQ ID NO: 4, (ii) a degenerate variant of SEQ ID NO: 4, or (iii) the complement of (i) or (ii), wherein the isolated nucleic acid is no more than about 100 kb in length, typically no more than about 75 kb in length, more typically no more than 5 about 50 kb in length. Often, the isolated nucleic acids of this embodiment are no more than about 25 kb in length, often no more than about 15 kb in length, and frequently no more than about 10 kb in length.

The invention also provides an isolated nucleic acid 10 comprising (i) a nucleotide sequence that encodes a peptide of at least 8 contiguous amino acids of SEQ ID NO: 5, (ii) a nucleotide sequence that encodes a peptide of at least 15 contiguous amino acids of SEQ ID NO: 5, or (iii) the complement of (i) or (ii), wherein the isolated nucleic acid is no more 15 than about 100 kb in length, typically no more than about 75 kb in length, more typically no more than about 50 kb in length. Often, the isolated nucleic acids of this embodiment are no more than about 25 kb in length, often no more than about 15 kb in length, and frequently no more than about 10 kb in length.

The invention also provides an isolated nucleic acid 20 comprising a nucleotide sequence that encodes (i) a polypeptide having the sequence of at least 8 contiguous amino acids of SEQ ID NO: 5 with conservative amino acid substitutions, (ii) a polypeptide having the sequence of at least 15 contiguous amino acids of SEQ ID NO: 5 with conservative amino acid substitutions, (iii) a polypeptide having the sequence of at least 8 contiguous amino acids of SEQ ID NO: 5 with moderately conservative substitutions, (iv) a polypeptide having the sequence of at least 15 contiguous amino acids of SEQ ID NO: 5 with moderately conservative substitutions, or (v) the complement of any of (i) - (iv), wherein the isolated nucleic acid is no more than about 100 kb in length, typically no more than about 75 kb in length, more typically no more than about 30

50 kb in length. Often, the isolated nucleic acids of this embodiment are no more than about 25 kb in length, often no more than about 15 kb in length, and frequently no more than about 10 kb in length.

5

Single Exon Probes

The invention further provides genome-derived single exon probes having portions of no more than one exon of the RGL3 gene. As further described in commonly owned and copending U.S. patent application serial no. 09/632,366, filed August 3, 2000 ("Methods and Apparatus for High Throughput Detection and Characterization of alternatively Spliced Genes"), the disclosure of which is incorporated herein by reference in its entirety, such single exon probes have particular utility in identifying and characterizing splice variants. In particular, such single exon probes are useful for identifying and discriminating the expression of distinct isoforms of RGL3.

In a first embodiment, the invention provides an isolated nucleic acid comprising a nucleotide sequence of no more than one portion of SEQ ID NOS: 6 - 24 or the complement of SEQ ID NOS: 6 - 24, wherein the portion comprises at least 17 contiguous nucleotides, 18 contiguous nucleotides, 20 contiguous nucleotides, 24 contiguous nucleotides, 25 contiguous nucleotides, or 50 contiguous nucleotides of any one of SEQ ID NOS: 6 - 24, or their complement. In a further embodiment, the exonic portion comprises the entirety of the referenced SEQ ID NO: or its complement.

In other embodiments, the invention provides isolated single exon probes having the nucleotide sequence of any one of SEQ ID NOS: 25 - 43.

Transcription Control Nucleic Acids

In another aspect, the present invention provides genome-derived isolated nucleic acids that include nucleic acid sequence elements that control transcription of the RGL3 gene.

5 These nucleic acids can be used, *inter alia*, to drive expression of heterologous coding regions in recombinant constructs, thus conferring upon such heterologous coding regions the expression pattern of the native RGL3 gene. These 10 nucleic acids can also be used, conversely, to target heterologous transcription control elements to the RGL3 genomic locus, altering the expression pattern of the RGL3 gene itself.

In a first such embodiment, the invention provides an isolated nucleic acid comprising the nucleotide sequence of SEQ 15 ID NO: 44 or its complement, wherein the isolated nucleic acid is no more than about 100 kb in length, typically no more than about 75 kb in length, more typically no more than about 50 kb in length. Often, the isolated nucleic acids of this embodiment are no more than about 25 kb in length, often no 20 more than about 15 kb in length, and frequently no more than about 10 kb in length.

In another embodiment, the invention provides an isolated nucleic acid comprising at least 17, 18, 20, 24, or 25 nucleotides of the sequence of SEQ ID NO: 44 or its complement, 25 wherein the isolated nucleic acid is no more than about 100 kb in length, typically no more than about 75 kb in length, more typically no more than about 50 kb in length. Often, the isolated nucleic acids of this embodiment are no more than about 25 kb in length, often no more than about 15 kb in 30 length, and frequently no more than about 10 kb in length.

VECTORS AND HOST CELLS

In another aspect, the present invention provides vectors that comprise one or more of the isolated nucleic acids of the present invention, and host cells in which such vectors have been introduced.

The vectors can be used, *inter alia*, for propagating the nucleic acids of the present invention in host cells (cloning vectors), for shuttling the nucleic acids of the present invention between host cells derived from disparate organisms (shuttle vectors), for inserting the nucleic acids of the present invention into host cell chromosomes (insertion vectors), for expressing sense or antisense RNA transcripts of the nucleic acids of the present invention *in vitro* or within a host cell, and for expressing polypeptides encoded by the nucleic acids of the present invention, alone or as fusions to heterologous polypeptides. Vectors of the present invention will often be suitable for several such uses.

Vectors are by now well-known in the art, and are described, *inter alia*, in Jones et al. (eds.), Vectors: Cloning Applications : Essential Techniques (Essential Techniques Series), John Wiley & Son Ltd 1998 (ISBN: 047196266X); Jones et al. (eds.), Vectors: Expression Systems : Essential Techniques (Essential Techniques Series), John Wiley & Son Ltd, 1998 (ISBN:0471962678); Gacesa et al., Vectors: Essential Data, John Wiley & Sons, 1995 (ISBN: 0471948411); Cid-Arregui (eds.), Viral Vectors: Basic Science and Gene Therapy, Eaton Publishing Co., 2000 (ISBN: 188129935X); Sambrook et al., Molecular Cloning: A Laboratory Manual (3rd ed.), Cold Spring Harbor Laboratory Press, 2001 (ISBN: 0879695773); Ausubel et al. (eds.), Short Protocols in Molecular Biology: A Compendium of Methods from Current Protocols in Molecular Biology (4th ed.), John Wiley & Sons, 1999 (ISBN: 047132938X), the disclosures of

which are incorporated herein by reference in their entireties. Furthermore, an enormous variety of vectors are available commercially. Use of existing vectors and modifications thereof being well within the skill in the art, only basic 5 features need be described here.

Typically, vectors are derived from virus, plasmid, prokaryotic or eukaryotic chromosomal elements, or some combination thereof, and include at least one origin of replication, at least one site for insertion of heterologous 10 nucleic acid, typically in the form of a polylinker with multiple, tightly clustered, single cutting restriction sites, and at least one selectable marker, although some integrative vectors will lack an origin that is functional in the host to be chromosomally modified, and some vectors will lack 15 selectable markers. Vectors of the present invention will further include at least one nucleic acid of the present invention inserted into the vector in at least one location.

Where present, the origin of replication and selectable markers are chosen based upon the desired host cell 20 or host cells; the host cells, in turn, are selected based upon the desired application.

For example, prokaryotic cells, typically *E. coli*, are typically chosen for cloning. In such case, vector replication is predicated on the replication strategies of 25 coliform-infecting phage - such as phage lambda, M13, T7, T3 and P1 - or on the replication origin of autonomously replicating episomes, notably the ColE1 plasmid and later derivatives, including pBR322 and the pUC series plasmids. Where *E. coli* is used as host, selectable markers are, 30 analogously, chosen for selectivity in gram negative bacteria: e.g., typical markers confer resistance to antibiotics, such as ampicillin, tetracycline, chloramphenicol, kanamycin, streptomycin, zeocin; auxotrophic markers can also be used.

As another example, yeast cells, typically *S. cerevisiae*, are chosen, *inter alia*, for eukaryotic genetic studies, due to the ease of targeting genetic changes by homologous recombination and to the ready ability to complement 5 genetic defects using recombinantly expressed proteins, for identification of interacting protein components, e.g. through use of a two-hybrid system, and for protein expression. Vectors of the present invention for use in yeast will typically, but not invariably, contain an origin of replication 10 suitable for use in yeast and a selectable marker that is functional in yeast.

Integrative YIp vectors do not replicate autonomously, but integrate, typically in single copy, into the yeast genome at low frequencies and thus replicate as part of 15 the host cell chromosome; these vectors lack an origin of replication that is functional in yeast, although they typically have at least one origin of replication suitable for propagation of the vector in bacterial cells. YEp vectors, in contrast, replicate episomally and autonomously due to presence 20 of the yeast 2 micron plasmid origin (2 μm ori). The YCp yeast centromere plasmid vectors are autonomously replicating vectors containing centromere sequences, CEN, and autonomously replicating sequences, ARS; the ARS sequences are believed to correspond to the natural replication origins of yeast 25 chromosomes. YACs are based on yeast linear plasmids, denoted YLp, containing homologous or heterologous DNA sequences that function as telomeres (TEL) *in vivo*, as well as containing yeast ARS (origins of replication) and CEN (centromeres) segments.

30 Selectable markers in yeast vectors include a variety of auxotrophic markers, the most common of which are (in *Saccharomyces cerevisiae*) URA3, HIS3, LEU2, TRP1 and LYS2, which complement specific auxotrophic mutations, such as

ura3-52, his3-D1, leu2-D1, trp1-D1 and lys2-201. The URA3 and LYS2 yeast genes further permit negative selection based on specific inhibitors, 5-fluoro-orotic acid (FOA) and α -amino adipic acid (α AA), respectively, that prevent growth of 5 the prototrophic strains but allows growth of the ura3 and lys2 mutants, respectively. Other selectable markers confer resistance to, e.g., zeocin.

As yet another example, insect cells are often chosen for high efficiency protein expression. Where the host cells 10 are from *Spodoptera frugiperda* – e.g., Sf9 and Sf21 cell lines, and expressSFTM cells (Protein Sciences Corp., Meriden, CT, USA) – the vector replicative strategy is typically based upon the baculovirus life cycle. Typically, baculovirus transfer vectors are used to replace the wild-type AcMNPV polyhedrin 15 gene with a heterologous gene of interest. Sequences that flank the polyhedrin gene in the wild-type genome are positioned 5' and 3' of the expression cassette on the transfer vectors. Following cotransfection with AcMNPV DNA, a homologous recombination event occurs between these sequences 20 resulting in a recombinant virus carrying the gene of interest and the polyhedrin or p10 promoter. Selection can be based upon visual screening for lacZ fusion activity.

As yet another example, mammalian cells are often chosen for expression of proteins intended as pharmaceutical 25 agents, and are also chosen as host cells for screening of potential agonist and antagonists of a protein or a physiological pathway.

Where mammalian cells are chosen as host cells, vectors intended for autonomous extrachromosomal replication 30 will typically include a viral origin, such as the SV40 origin (for replication in cell lines expressing the large T-antigen, such as COS1 and COS7 cells), the papillomavirus origin, or the EBV origin for long term episomal replication (for use, e.g.,

in 293-EBNA cells, which constitutively express the EBV EBNA-1 gene product and adenovirus E1A). Vectors intended for integration, and thus replication as part of the mammalian chromosome, can, but need not, include an origin of replication functional in mammalian cells, such as the SV40 origin. 5 Vectors based upon viruses, such as adenovirus, adeno-associated virus, vaccinia virus, and various mammalian retroviruses, will typically replicate according to the viral replicative strategy.

10 Selectable markers for use in mammalian cells include resistance to neomycin (G418), blasticidin, hygromycin and to zeocin, and selection based upon the purine salvage pathway using HAT medium.

15 Plant cells can also be used for expression, with the vector replicon typically derived from a plant virus (e.g., cauliflower mosaic virus, CaMV; tobacco mosaic virus, TMV) and selectable markers chosen for suitability in plants.

20 For propagation of nucleic acids of the present invention that are larger than can readily be accommodated in vectors derived from plasmids or virus, the invention further provides artificial chromosomes – BACs, YACs, PACs, and HACs – that comprise RGL3 nucleic acids, often genomic nucleic acids.

25 The BAC system is based on the well-characterized *E. coli* F-factor, a low copy plasmid that exists in a supercoiled circular form in host cells. The structural features of the F-factor allow stable maintenance of individual human DNA clones as well as easy manipulation of the cloned DNA. See Shizuya et al., *Keio J. Med.* 50(1):26-30 (2001); Shizuya et al., *Proc. Natl. Acad. Sci. USA* 89(18):8794-7 (1992).

30 YACs are based on yeast linear plasmids, denoted YLp, containing homologous or heterologous DNA sequences that function as telomeres (TEL) *in vivo*, as well as containing

yeast ARS (origins of replication) and CEN (centromeres) segments.

HACs are human artificial chromosomes. Kuroiwa et al., *Nature Biotechnol.* 18(10):1086-90 (2000); Henning et al., 5 *Proc. Natl. Acad. Sci. USA* 96(2):592-7 (1999); Harrington et al., *Nature Genet.* 15(4):345-55 (1997). In one version, long synthetic arrays of alpha satellite DNA are combined with telomeric DNA and genomic DNA to generate linear microchromosomes that are mitotically and cytogenetically 10 stable in the absence of selection.

PACs are P1-derived artificial chromosomes. Sternberg, *Proc. Natl. Acad. Sci. USA* 87(1):103-7 (1990); Sternberg et al., *New Biol.* 2(2):151-62 (1990); Pierce et al., *Proc. Natl Acad. Sci. USA* 89(6):2056-60 (1992).

15 Vectors of the present invention will also often include elements that permit *in vitro* transcription of RNA from the inserted heterologous nucleic acid. Such vectors typically include a phage promoter, such as that from T7, T3, or SP6, flanking the nucleic acid insert. Often two different 20 such promoters flank the inserted nucleic acid, permitting separate *in vitro* production of both sense and antisense strands.

Expression vectors of the present invention – that is, those vectors that will drive expression of polypeptides 25 from the inserted heterologous nucleic acid – will often include a variety of other genetic elements operatively linked to the protein-encoding heterologous nucleic acid insert, typically genetic elements that drive transcription, such as promoters and enhancer elements, those that facilitate RNA 30 processing, such as transcription termination and/or polyadenylation signals, and those that facilitate translation, such as ribosomal consensus sequences.

For example, vectors for expressing proteins of the present invention in prokaryotic cells, typically *E. coli*, will include a promoter, often a phage promoter, such as phage lambda pL promoter, the trc promoter, a hybrid derived from the 5 trp and lac promoters, the bacteriophage T7 promoter (in *E. coli* cells engineered to express the T7 polymerase), or the araBAD operon. Often, such prokaryotic expression vectors will further include transcription terminators, such as the aspA 10 terminator, and elements that facilitate translation, such as a consensus ribosome binding site and translation termination codon, Schomer et al., *Proc. Natl. Acad. Sci. USA* 83:8506-8510 (1986).

As another example, vectors for expressing proteins of the present invention in yeast cells, typically *S. cerevisiae*, will include a yeast promoter, such as the CYC1 15 promoter, the GAL1 promoter, ADH1 promoter, or the GPD promoter, and will typically have elements that facilitate transcription termination, such as the transcription termination signals from the CYC1 or ADH1 gene.

As another example, vectors for expressing proteins of the present invention in mammalian cells will include a promoter active in mammalian cells. Such promoters are often drawn from mammalian viruses – such as the enhancer-promoter sequences from the immediate early gene of the human 20 cytomegalovirus (CMV), the enhancer-promoter sequences from the Rous sarcoma virus long terminal repeat (RSV LTR), and the enhancer-promoter from SV40. Often, expression is enhanced by incorporation of polyadenylation sites, such as the late SV40 25 polyadenylation site and the polyadenylation signal and transcription termination sequences from the bovine growth hormone (BGH) gene, and ribosome binding sites. Furthermore, 30 vectors can include introns, such as intron II of rabbit β-globin gene and the SV40 splice elements.

Vector-drive protein expression can be constitutive or inducible.

Inducible vectors include either naturally inducible promoters, such as the trc promoter, which is regulated by the lac operon, and the pL promoter, which is regulated by tryptophan, the MMTV-LTR promoter, which is inducible by dexamethasone, or can contain synthetic promoters and/or additional elements that confer inducible control on adjacent promoters. Examples of inducible synthetic promoters are the hybrid Plac/ara-1 promoter and the PLtetO-1 promoter. The PLtetO-1 promoter takes advantage of the high expression levels from the PL promoter of phage lambda, but replaces the lambda repressor sites with two copies of operator 2 of the Tn10 tetracycline resistance operon, causing this promoter to be tightly repressed by the Tet repressor protein and induced in response to tetracycline (Tc) and Tc derivatives such as anhydrotetracycline.

As another example of inducible elements, hormone response elements, such as the glucocorticoid response element (GRE) and the estrogen response element (ERE), can confer hormone inducibility where vectors are used for expression in cells having the respective hormone receptors. To reduce background levels of expression, elements responsive to ecdysone, an insect hormone, can be used instead, with coexpression of the ecdysone receptor.

Expression vectors can be designed to fuse the expressed polypeptide to small protein tags that facilitate purification and/or visualization.

For example, proteins of the present invention can be expressed with a polyhistidine tag that facilitates purification of the fusion protein by immobilized metal affinity chromatography, for example using NiNTA resin (Qiagen Inc., Valencia, CA, USA) or TALONTM resin (cobalt immobilized

affinity chromatography medium, Clontech Labs, Palo Alto, CA, USA). As another example, the fusion protein can include a chitin-binding tag and self-excising intein, permitting chitin-based purification with self-removal of the fused tag (IMPACTTM system, New England Biolabs, Inc., Beverley, MA, USA). Alternatively, the fusion protein can include a calmodulin-binding peptide tag, permitting purification by calmodulin affinity resin (Stratagene, La Jolla, CA, USA), or a specifically excisable fragment of the biotin carboxylase carrier protein, permitting purification of *in vivo* biotinylated protein using an avidin resin and subsequent tag removal (Promega, Madison, WI, USA). As another useful alternative, the proteins of the present invention can be expressed as a fusion to glutathione-S-transferase, the affinity and specificity of binding to glutathione permitting purification using glutathione affinity resins, such as Glutathione-Superflow Resin (Clontech Laboratories, Palo Alto, CA, USA), with subsequent elution with free glutathione.

Other tags include, for example, the Xpress epitope, detectable by anti-Xpress antibody (Invitrogen, Carlsbad, CA, USA), a myc tag, detectable by anti-myc tag antibody, the V5 epitope, detectable by anti-V5 antibody (Invitrogen, Carlsbad, CA, USA), FLAG[®] epitope, detectable by anti-FLAG[®] antibody (Stratagene, La Jolla, CA, USA), and the HA epitope.

For secretion of expressed proteins, vectors can include appropriate sequences that encode secretion signals, such as leader peptides. For example, the pSecTag2 vectors (Invitrogen, Carlsbad, CA, USA) are 5.2 kb mammalian expression vectors that carry the secretion signal from the V-J2-C region of the mouse Ig kappa-chain for efficient secretion of recombinant proteins from a variety of mammalian cell lines.

Expression vectors can also be designed to fuse proteins encoded by the heterologous nucleic acid insert to

polypeptides larger than purification and/or identification tags. Useful protein fusions include those that permit display of the encoded protein on the surface of a phage or cell, fusions to intrinsically fluorescent proteins, such as those 5 that have a green fluorescent protein (GFP)-like chromophore, fusions to the IgG Fc region, and fusions for use in two hybrid systems.

Vectors for phage display fuse the encoded polypeptide to, e.g., the gene III protein (pIII) or gene VIII 10 protein (pVIII) for display on the surface of filamentous phage, such as M13. See Barbas et al., Phage Display: A Laboratory Manual, Cold Spring Harbor Laboratory Press (2001) (ISBN 0-87969-546-3); Kay et al. (eds.), Phage Display of Peptides and Proteins: A Laboratory Manual, San Diego: Academic 15 Press, Inc., 1996; Abelson et al. (eds.), Combinatorial Chemistry, Methods in Enzymology vol. 267, Academic Press (May 1996).

Vectors for yeast display, e.g. the pYD1 yeast display vector (Invitrogen, Carlsbad, CA, USA), use the 20 α -agglutinin yeast adhesion receptor to display recombinant protein on the surface of *S. cerevisiae*. Vectors for mammalian display, e.g., the pDisplay™ vector (Invitrogen, Carlsbad, CA, USA), target recombinant proteins using an N-terminal cell surface targeting signal and a C-terminal transmembrane 25 anchoring domain of platelet derived growth factor receptor.

A wide variety of vectors now exist that fuse proteins encoded by heterologous nucleic acids to the chromophore of the substrate-independent, intrinsically 30 fluorescent green fluorescent protein from *Aequorea victoria* ("GFP") and its variants. These proteins are intrinsically fluorescent: the GFP-like chromophore is entirely encoded by its amino acid sequence and can fluoresce without requirement for cofactor or substrate.

Structurally, the GFP-like chromophore comprises an 11-stranded β -barrel (β -can) with a central α -helix, the central α -helix having a conjugated π -resonance system that includes two aromatic ring systems and the bridge between them.

5 The π -resonance system is created by autocatalytic cyclization among amino acids; cyclization proceeds through an imidazolinone intermediate, with subsequent dehydrogenation by molecular oxygen at the $\text{C}\alpha\text{-C}\beta$ bond of a participating tyrosine.

The GFP-like chromophore can be selected from GFP-like chromophores found in naturally occurring proteins, such as *A. victoria* GFP (GenBank accession number AAA27721), *Renilla reniformis* GFP, FP583 (GenBank accession no. AF168419) (DsRed), FP593 (AF272711), FP483 (AF168420), FP484 (AF168424), FP595 (AF246709), FP486 (AF168421), FP538 (AF168423), and FP506 (AF168422), and need include only so much of the native protein as is needed to retain the chromophore's intrinsic fluorescence. Methods for determining the minimal domain required for fluorescence are known in the art. Li et al., "Deletions of the *Aequorea victoria* Green Fluorescent Protein Define the Minimal Domain Required for Fluorescence," *J. Biol. Chem.* 272:28545-28549 (1997).

Alternatively, the GFP-like chromophore can be selected from GFP-like chromophores modified from those found in nature. Typically, such modifications are made to improve recombinant production in heterologous expression systems (with or without change in protein sequence), to alter the excitation and/or emission spectra of the native protein, to facilitate purification, to facilitate or as a consequence of cloning, or are a fortuitous consequence of research investigation.

30 The methods for engineering such modified GFP-like chromophores and testing them for fluorescence activity, both alone and as part of protein fusions, are well-known in the art. Early results of these efforts are reviewed in Heim et

al., *Curr. Biol.* 6:178-182 (1996), incorporated herein by reference in its entirety; a more recent review, with tabulation of useful mutations, is found in Palm et al., "Spectral Variants of Green Fluorescent Protein," in Green Fluorescent Proteins, Conn (ed.), *Methods Enzymol.* vol. 302, 5 pp. 378 - 394 (1999), incorporated herein by reference in its entirety. A variety of such modified chromophores are now commercially available and can readily be used in the fusion proteins of the present invention.

For example, EGFP ("enhanced GFP"), Cormack et al., *Gene* 173:33-38 (1996); U.S. Pat. Nos. 6,090,919 and 5,804,387, is a red-shifted, human codon-optimized variant of GFP that has been engineered for brighter fluorescence, higher expression in mammalian cells, and for an excitation spectrum optimized for 10 use in flow cytometers. EGFP can usefully contribute a GFP-like chromophore to the fusion proteins of the present invention. A variety of EGFP vectors, both plasmid and viral, are available commercially (Clontech Labs, Palo Alto, CA, USA), including vectors for bacterial expression, vectors for N-terminal protein fusion expression, vectors for expression of C-terminal protein fusions, and for bicistronic expression. 15 20

Toward the other end of the emission spectrum, EBFP ("enhanced blue fluorescent protein") and BFP2 contain four amino acid substitutions that shift the emission from green to 25 blue, enhance the brightness of fluorescence and improve solubility of the protein, Heim et al., *Curr. Biol.* 6:178-182 (1996); Cormack et al., *Gene* 173:33-38 (1996). EBFP is optimized for expression in mammalian cells whereas BFP2, which retains the original jellyfish codons, can be expressed in 30 bacteria; as is further discussed below, the host cell of production does not affect the utility of the resulting fusion protein. The GFP-like chromophores from EBFP and BFP2 can usefully be included in the fusion proteins of the present

invention, and vectors containing these blue-shifted variants are available from Clontech Labs (Palo Alto, CA, USA).

Analogously, EYFP ("enhanced yellow fluorescent protein"), also available from Clontech Labs, contains four 5 amino acid substitutions, different from EBFP, Ormö et al., *Science* 273:1392-1395 (1996), that shift the emission from green to yellowish-green. Citrine, an improved yellow fluorescent protein mutant, is described in Heikal et al., *Proc. Natl. Acad. Sci. USA* 97:11996-12001 (2000). ECFP 10 ("enhanced cyan fluorescent protein") (Clontech Labs, Palo Alto, CA, USA) contains six amino acid substitutions, one of which shifts the emission spectrum from green to cyan. Heim et al., *Curr. Biol.* 6:178-182 (1996); Miyawaki et al., *Nature* 388:882-887 (1997). The GFP-like chromophore of each of these 15 GFP variants can usefully be included in the fusion proteins of the present invention.

The GFP-like chromophore can also be drawn from other modified GFPs, including those described in U.S. Pat. Nos. 6,124,128; 6,096,865; 6,090,919; 6,066,476; 6,054,321; 20 6,027,881; 5,968,750; 5,874,304; 5,804,387; 5,777,079; 5,741,668; and 5,625,048, the disclosures of which are incorporated herein by reference in their entireties. See also Conn (ed.), Green Fluorescent Protein, Methods in Enzymol. Vol. 302, pp 378-394 (1999), incorporated herein by reference in its entirety. A variety of such modified chromophores are now 25 commercially available and can readily be used in the fusion proteins of the present invention.

Fusions to the IgG Fc region increase serum half life of protein pharmaceutical products through interaction with the 30 FcRn receptor (also denominated the FcRp receptor and the Brambell receptor, FcRb), further described in international patent application nos. WO 97/43316, WO 97/34631, WO 96/32478, WO 96/18412.

For long-term, high-yield recombinant production of the proteins, protein fusions, and protein fragments of the present invention, stable expression is particularly useful.

Stable expression is readily achieved by integration 5 into the host cell genome of vectors having selectable markers, followed by selection for integrants.

For example, the pUB6/V5-His A, B, and C vectors (Invitrogen, Carlsbad, CA, USA) are designed for high-level stable expression of heterologous proteins in a wide range of 10 mammalian tissue types and cell lines. pUB6/V5-His uses the promoter/enhancer sequence from the human ubiquitin C gene to drive expression of recombinant proteins: expression levels in 293, CHO, and NIH3T3 cells are comparable to levels from the CMV and human EF-1 α promoters. The bsd gene permits rapid 15 selection of stably transfected mammalian cells with the potent antibiotic blasticidin.

Replication incompetent retroviral vectors, typically derived from Moloney murine leukemia virus, prove particularly useful for creating stable transfectants having integrated 20 provirus. The highly efficient transduction machinery of retroviruses, coupled with the availability of a variety of packaging cell lines – such as RetroPackTM PT 67, EcoPack2TM – 293, AmphotroPack-293, GP2-293 cell lines (all available from Clontech Laboratories, Palo Alto, CA, USA) – allow a wide host 25 range to be infected with high efficiency; varying the multiplicity of infection readily adjusts the copy number of the integrated provirus. Retroviral vectors are available with a variety of selectable markers, such as resistance to neomycin, hygromycin, and puromycin, permitting ready selection 30 of stable integrants.

The present invention further includes host cells comprising the vectors of the present invention, either present

episomally within the cell or integrated, in whole or in part, into the host cell chromosome.

Among other considerations, some of which are described above, a host cell strain may be chosen for its ability to process the expressed protein in the desired fashion. Such post-translational modifications of the polypeptide include, but are not limited to, acetylation, carboxylation, glycosylation, phosphorylation, lipidation, and acylation, and it is an aspect of the present invention to provide RGL3 proteins with such post-translational modifications.

As noted earlier, host cells can be prokaryotic or eukaryotic. Representative examples of appropriate host cells include, but are not limited to, bacterial cells, such as *E. coli*, *Caulobacter crescentus*, *Streptomyces* species, and *Salmonella typhimurium*; yeast cells, such as *Saccharomyces cerevisiae*, *Schizosaccharomyces pombe*, *Pichia pastoris*, *Pichia methanolica*; insect cell lines, such as those from *Spodoptera frugiperda* - e.g., Sf9 and Sf21 cell lines, and expressSFTM cells (Protein Sciences Corp., Meriden, CT, USA) - *Drosophila S2* cells, and *Trichoplusia ni* High Five[®] Cells (Invitrogen, Carlsbad, CA, USA); and mammalian cells. Typical mammalian cells include COS1 and COS7 cells, Chinese hamster ovary (CHO) cells, NIH 3T3 cells, 293 cells, HEPG2 cells, HeLa cells, L cells, murine ES cell lines (e.g., from strains 129/SV, C57/BL6, DBA-1, 129/SVJ), K562, Jurkat cells, and BW5147. Other mammalian cell lines are well known and readily available from the American Type Culture Collection (ATCC) (Manassas, VA, USA) and the National Institute of General Medical Sciences (NIGMS) Human Genetic Cell Repository at the Coriell Cell Repositories (Camden, NJ, USA).

Methods for introducing the vectors and nucleic acids of the present invention into the host cells are well known in

the art; the choice of technique will depend primarily upon the specific vector to be introduced and the host cell chosen.

For example, phage lambda vectors will typically be packaged using a packaging extract (e.g., Gigapack® packaging extract, Stratagene, La Jolla, CA, USA), and the packaged virus used to infect *E. coli*. Plasmid vectors will typically be introduced into chemically competent or electrocompetent bacterial cells.

E. coli cells can be rendered chemically competent by treatment, e.g., with CaCl₂, or a solution of Mg²⁺, Mn²⁺, Ca²⁺, Rb⁺ or K⁺, dimethyl sulfoxide, dithiothreitol, and hexamine cobalt (III), Hanahan, *J. Mol. Biol.* 166(4):557-80 (1983), and vectors introduced by heat shock. A wide variety of chemically competent strains are also available commercially (e.g., Epicurian Coli® XL10-Gold® Ultracompetent Cells (Stratagene, La Jolla, CA, USA); DH5α competent cells (Clontech Laboratories, Palo Alto, CA, USA); TOP10 Chemically Competent *E. coli* Kit (Invitrogen, Carlsbad, CA, USA)).

Bacterial cells can be rendered electrocompetent – that is, competent to take up exogenous DNA by electroporation – by various pre-pulse treatments; vectors are introduced by electroporation followed by subsequent outgrowth in selected media. An extensive series of protocols is provided online in Electroprotocols (BioRad, Richmond, CA, USA) (http://www.bio-rad.com/LifeScience/pdf/New_Gene_Pulser.pdf).

Vectors can be introduced into yeast cells by spheroplasting, treatment with lithium salts, electroporation, or protoplast fusion.

Spheroplasts are prepared by the action of hydrolytic enzymes – a snail-gut extract, usually denoted Glusulase, or Zymolyase, an enzyme from *Arthrobacter luteus* – to remove portions of the cell wall in the presence of osmotic stabilizers, typically 1 M sorbitol. DNA is added to the

spheroplasts, and the mixture is co-precipitated with a solution of polyethylene glycol (PEG) and Ca^{2+} . Subsequently, the cells are resuspended in a solution of sorbitol, mixed with molten agar and then layered on the surface of a selective plate containing sorbitol. For lithium-mediated transformation, yeast cells are treated with lithium acetate, which apparently permeabilizes the cell wall, DNA is added and the cells are co-precipitated with PEG. The cells are exposed to a brief heat shock, washed free of PEG and lithium acetate, and subsequently spread on plates containing ordinary selective medium. Increased frequencies of transformation are obtained by using specially-prepared single-stranded carrier DNA and certain organic solvents. Schiestl et al., *Curr. Genet.* 16(5-6):339-46 (1989). For electroporation, freshly-grown yeast cultures are typically washed, suspended in an osmotic protectant, such as sorbitol, mixed with DNA, and the cell suspension pulsed in an electroporation device. Subsequently, the cells are spread on the surface of plates containing selective media. Becker et al., *Methods Enzymol.* 194:182-7 (1991). The efficiency of transformation by electroporation can be increased over 100-fold by using PEG, single-stranded carrier DNA and cells that are in late log-phase of growth. Larger constructs, such as YACs, can be introduced by protoplast fusion.

Mammalian and insect cells can be directly infected by packaged viral vectors, or transfected by chemical or electrical means.

For chemical transfection, DNA can be coprecipitated with CaPO_4 or introduced using liposomal and nonliposomal lipid-based agents. Commercial kits are available for CaPO_4 transfection (CalPhos™ Mammalian Transfection Kit, Clontech Laboratories, Palo Alto, CA, USA), and lipid-mediated transfection can be practiced using commercial reagents, such

- as LIPOFECTAMINE™ 2000, LIPOFECTAMINE™ Reagent, CELLFECTIN® Reagent, and LIPOFECTIN® Reagent (Invitrogen, Carlsbad, CA, USA), DOTAP Liposomal Transfection Reagent, FuGENE 6, X-tremeGENE Q2, DOSPER, (Roche Molecular Biochemicals, Indianapolis, IN USA), Effectene™, PolyFect®, Superfect® (Qiagen, Inc., Valencia, CA, USA). Protocols for electroporating mammalian cells can be found online in Electroprotocols (Bio-Rad, Richmond, CA, USA) (http://www.bio-rad.com/LifeScience/pdf/New_Gene_Pulser.pdf).
- 5 See also, Norton et al. (eds.), Gene Transfer Methods: Introducing DNA into Living Cells and Organisms, BioTechniques Books, Eaton Publishing Co. (2000) (ISBN 1-881299-34-1), incorporated herein by reference in its entirety.
- 10 Other transfection techniques include transfection by particle embardment. See, e.g., Cheng et al., *Proc. Natl. Acad. Sci. USA* 90(10):4455-9 (1993); Yang et al., *Proc. Natl. Acad. Sci. USA* 87(24):9568-72 (1990).

PROTEINS

- 20 In another aspect, the present invention provides RGL3 proteins, various fragments thereof suitable for use as antigens (e.g., for epitope mapping) and for use as immunogens (e.g., for raising antibodies or as vaccines), fusions of RGL3 polypeptides and fragments to heterologous polypeptides, and conjugates of the proteins, fragments, and fusions of the present invention to other moieties (e.g., to carrier proteins, to fluorophores).

25 FIG. 3 presents the predicted amino acid sequences encoded by the RGL3 cDNA clone. The amino acid sequence is further presented, respectively, in SEQ ID NO: 3.

30 Unless otherwise indicated, amino acid sequences of the proteins of the present invention were determined as a

predicted translation from a nucleic acid sequence. Accordingly, any amino acid sequence presented herein may contain errors due to errors in the nucleic acid sequence, as described in detail above. Furthermore, single nucleotide polymorphisms (SNPs) occur frequently in eukaryotic genomes - more than 1.4 million SNPs have already identified in the human genome, International Human Genome Sequencing Consortium, *Nature* 409:860 - 921 (2001) - and the sequence determined from one individual of a species may differ from other allelic forms present within the population. Small deletions and insertions can often be found that do not alter the function of the protein.

Accordingly, it is an aspect of the present invention to provide proteins not only identical in sequence to those described with particularity herein, but also to provide isolated proteins at least about 65% identical in sequence to those described with particularity herein, typically at least about 70%, 75%, 80%, 85%, or 90% identical in sequence to those described with particularity herein, usefully at least about 91%, 92%, 93%, 94%, or 95% identical in sequence to those described with particularity herein, usefully at least about 96%, 97%, 98%, or 99% identical in sequence to those described with particularity herein, and, most conservatively, at least about 99.5%, 99.6%, 99.7%, 99.8% and 99.9% identical in sequence to those described with particularity herein. These sequence variants can be naturally occurring or can result from human intervention by way of random or directed mutagenesis.

For purposes herein, percent identity of two amino acid sequences is determined using the procedure of Tatiana et al., "Blast 2 sequences - a new tool for comparing protein and nucleotide sequences", *FEMS Microbiol Lett.* 174:247-250 (1999), which procedure is effectuated by the computer program BLAST 2 SEQUENCES, available online at

<http://www.ncbi.nlm.nih.gov/blast/bl2seq/bl2.html>,

To assess percent identity of amino acid sequences, the BLASTP
 5 module of BLAST 2 SEQUENCES is used with default values of (i)
 BLOSUM62 matrix, Henikoff et al., *Proc. Natl. Acad. Sci USA*
 89(22):10915-9 (1992); (ii) open gap 11 and extension gap 1
 penalties; and (iii) gap x_dropoff 50 expect 10 word size 3
 filter, and both sequences are entered in their entireties.

10 As is well known, amino acid substitutions occur
 frequently among natural allelic variants, with conservative
 substitutions often occasioning only *de minimis* change in
 protein function.

Accordingly, it is an aspect of the present invention
 15 to provide proteins not only identical in sequence to those
 described with particularity herein, but also to provide
 isolated proteins having the sequence of RGL3 proteins, or
 portions thereof, with conservative amino acid substitutions.
 It is a further aspect to provide isolated proteins having the
 20 sequence of RGL3 proteins, and portions thereof, with
 moderately conservative amino acid substitutions. These
 conservatively-substituted and moderately conservatively-
 substituted variants can be naturally occurring or can result
 from human intervention.

25 Although there are a variety of metrics for calling
 conservative amino acid substitutions, based primarily on
 either observed changes among evolutionarily related proteins
 or on predicted chemical similarity, for purposes herein a
 conservative replacement is any change having a positive value
 30 in the PAM250 log-likelihood matrix reproduced herein below
 (see Gonnet et al., *Science* 256(5062):1443-5 (1992)):

	A	R	N	D	C	Q	E	G	H	I	L	K	M	F	P	S	T	W	Y	V
A	2	-1	0	0	0	0	0	0	-1	-1	-1	0	-1	-2	0	1	1	-4	-2	0

R	-1	5	0	0	-2	2	0	-1	1	-2	-2	3	-2	-3	-1	0	0	-2	-2	-2
N	0	0	4	2	-2	1	1	0	1	-3	-3	1	-2	-3	-1	1	0	-4	-1	-2
D	0	0	2	5	-3	1	3	0	0	-4	-4	0	-3	-4	-1	0	0	-5	-3	-3
C	0	-2	-2	-3	12	-2	-3	-2	-1	-1	-2	-3	-1	-1	-3	0	0	-1	0	0
5	Q	0	2	1	1	-2	3	2	-1	1	-2	-2	2	-1	-3	0	0	0	-3	-2
E	0	0	1	3	-3	2	4	-1	0	-3	-3	1	-2	-4	0	0	0	-4	-3	-2
G	0	-1	0	0	-2	-1	-1	7	-1	-4	-4	-1	-4	-5	-2	0	-1	-4	-4	-3
H	-1	1	1	0	-1	1	0	-1	6	-2	-2	1	-1	0	-1	0	0	-1	2	-2
I	-1	-2	-3	-4	-1	-2	-3	-4	-2	4	3	-2	2	1	-3	-2	-1	-2	-1	3
10	L	-1	-2	-3	-4	-2	-2	-3	-4	-2	3	4	-2	3	2	-2	-2	-1	-1	0
K	0	3	1	0	-3	2	1	-1	1	-2	-2	3	-1	-3	-1	0	0	-4	-2	-2
M	-1	-2	-2	-3	-1	-1	-2	-4	-1	2	3	-1	4	2	-2	-1	-1	-1	0	2
F	-2	-3	-3	-4	-1	-3	-4	-5	0	1	2	-3	2	7	-4	-3	-2	4	5	0
P	0	-1	-1	-1	-3	0	0	-2	-1	-3	-2	-1	-2	-4	8	0	0	-5	-3	-2
15	S	1	0	1	0	0	0	0	0	-2	-2	0	-1	-3	0	2	2	-3	-2	-1
T	1	0	0	0	0	0	0	-1	0	-1	-1	0	-1	-2	0	2	2	-4	-2	0
W	-4	-2	-4	-5	-1	-3	-4	-4	-1	-2	-1	-4	-1	4	-5	-3	-4	14	4	-3
Y	-2	-2	-1	-3	0	-2	-3	-4	2	-1	0	-2	0	5	-3	-2	-2	4	8	-1
V	0	-2	-2	-3	0	-2	-2	-3	-2	3	2	-2	2	0	-2	-1	0	-3	-1	3

20

For purposes herein, a "moderately conservative" replacement is any change having a nonnegative value in the PAM250 log-likelihood matrix reproduced herein above.

As is also well known in the art, relatedness of proteins can also be characterized using a functional test, the ability of the encoding nucleic acids to base-pair to one another at defined hybridization stringencies.

It is, therefore, another aspect of the invention to provide isolated proteins not only identical in sequence to those described with particularity herein, but also to provide isolated proteins ("hybridization related proteins") that are encoded by nucleic acids that hybridize under high stringency conditions (as defined herein above) to all or to a portion of various of the isolated nucleic acids of the present invention ("reference nucleic acids"). It is a further aspect of the invention to provide isolated proteins ("hybridization related

proteins") that are encoded by nucleic acids that hybridize under moderate stringency conditions (as defined herein above) to all or to a portion of various of the isolated nucleic acids of the present invention ("reference nucleic acids").

5 The hybridization related proteins can be alternative isoforms, homologues, paralogues, and orthologues of the RGL3 protein of the present invention. Particularly useful orthologues are those from other primate species, such as chimpanzee, rhesus macaque monkey, baboon, orangutan, and 10 gorilla, from rodents, such as rats, mice, guinea pigs; from lagomorphs, such as rabbits, and from domestic livestock, such as cow, pig, sheep, horse, and goat.

15 Relatedness of proteins can also be characterized using a second functional test, the ability of a first protein competitively to inhibit the binding of a second protein to an antibody.

20 It is, therefore, another aspect of the present invention to provide isolated proteins not only identical in sequence to those described with particularity herein, but also to provide isolated proteins ("cross-reactive proteins") that competitively inhibit the binding of antibodies to all or to a portion of various of the isolated RGL3 proteins of the present invention ("reference proteins"). Such competitive inhibition can readily be determined using immunoassays well known in the 25 art.

30 Among the proteins of the present invention that differ in amino acid sequence from those described with particularity herein – including those that have deletions and insertions causing up to 10% non-identity, those having conservative or moderately conservative substitutions, hybridization related proteins, and cross-reactive proteins – those that substantially retain one or more RGL3 activities are particularly useful. As described above, those activities

include guanine nucleotide exchange for the small GTPase Ral and being a downstream effector for both Rit and Ras.

Residues that are tolerant of change while retaining function can be identified by altering the protein at known residues using methods known in the art, such as alanine scanning mutagenesis, Cunningham et al., *Science* 244(4908):1081-5 (1989); transposon linker scanning mutagenesis, Chen et al., *Gene* 263(1-2):39-48 (2001); combinations of homolog- and alanine-scanning mutagenesis, Jin et al., *J. Mol. Biol.* 226(3):851-65 (1992); combinatorial alanine scanning, Weiss et al., *Proc. Natl. Acad. Sci USA* 97(16):8950-4 (2000), followed by functional assay. Transposon linker scanning kits are available commercially (New England Biolabs, Beverly, MA, USA, catalog. no. E7-102S; EZ::TN™ 15 In-Frame Linker Insertion Kit, catalogue no. EZI04KN, Epicentre Technologies Corporation, Madison, WI, USA).

As further described below, the isolated proteins of the present invention can readily be used as specific immunogens to raise antibodies that specifically recognize RGL3 proteins, their isoforms, homologues, paralogues, and/or orthologues. The antibodies, in turn, can be used, *inter alia*, specifically to assay for the RGL3 proteins of the present invention – e.g. by ELISA for detection of protein fluid samples, such as serum, by immunohistochemistry or laser scanning cytometry, for detection of protein in tissue samples, or by flow cytometry, for detection of intracellular protein in cell suspensions – for specific antibody-mediated isolation and/or purification of RGL3 proteins, as for example by immunoprecipitation, and for use as specific agonists or antagonists of RGL3 action.

The isolated proteins of the present invention are also immediately available for use as specific standards in assays used to determine the concentration and/or amount

specifically of the RGL3 proteins of the present invention. As is well known, ELISA kits for detection and quantitation of protein analytes typically include isolated and purified protein of known concentration for use as a measurement standard (e.g., the human interferon- γ OptEIA kit, catalog no. 5 555142, Pharmingen, San Diego, CA, USA includes human recombinant gamma interferon, baculovirus produced).

The isolated proteins of the present invention are also immediately available for use as specific biomolecule 10 capture probes for surface-enhanced laser desorption ionization (SELDI) detection of protein-protein interactions, WO 98/59362; WO 98/59360; WO 98/59361; and Merchant *et al.*, *Electrophoresis* 21(6):1164-77 (2000), the disclosures of which are incorporated herein by reference in their entireties. Analogously, the 15 isolated proteins of the present invention are also immediately available for use as specific biomolecule capture probes on BIACORE surface plasmon resonance probes. . See Weinberger *et al.*, *Pharmacogenomics* 1(4):395-416 (2000); Malmqvist, *Biochem. Soc. Trans.* 27(2):335-40 (1999).

20 The isolated proteins of the present invention are also useful as a therapeutic supplement in patients having a specific deficiency in RGL3 production.

In another aspect, the invention also provides fragments of various of the proteins of the present invention. 25 The protein fragments are useful, *inter alia*, as antigenic and immunogenic fragments of RGL3.

By "fragments" of a protein is here intended isolated proteins (equally, polypeptides, peptides, oligopeptides), however obtained, that have an amino acid sequence identical to 30 a portion of the reference amino acid sequence, which portion is at least 6 amino acids and less than the entirety of the reference nucleic acid. As so defined, "fragments" need not be

obtained by physical fragmentation of the reference protein, although such provenance is not thereby precluded.

Fragments of at least 6 contiguous amino acids are useful in mapping B cell and T cell epitopes of the reference protein. See, e.g., Geysen et al., "Use of peptide synthesis to probe viral antigens for epitopes to a resolution of a single amino acid," *Proc. Natl. Acad. Sci. USA* 81:3998-4002 (1984) and U.S. Pat. Nos. 4,708,871 and 5,595,915, the disclosures of which are incorporated herein by reference in their entireties. Because the fragment need not itself be immunogenic, part of an immunodominant epitope, nor even recognized by native antibody, to be useful in such epitope mapping, all fragments of at least 6 amino acids of the proteins of the present invention have utility in such a study.

Fragments of at least 8 contiguous amino acids, often at least 15 contiguous amino acids, have utility as immunogens for raising antibodies that recognize the proteins of the present invention. See, e.g., Lerner, "Tapping the immunological repertoire to produce antibodies of predetermined specificity," *Nature* 299:592-596 (1982); Shinnick et al., "Synthetic peptide immunogens as vaccines," *Annu. Rev. Microbiol.* 37:425-46 (1983); Sutcliffe et al., "Antibodies that react with predetermined sites on proteins," *Science* 219:660-6 (1983), the disclosures of which are incorporated herein by reference in their entireties. As further described in the above-cited references, virtually all 8-mers, conjugated to a carrier, such as a protein, prove immunogenic - that is, prove capable of eliciting antibody for the conjugated peptide; accordingly, all fragments of at least 8 amino acids of the proteins of the present invention have utility as immunogens.

Fragments of at least 8, 9, 10 or 12 contiguous amino acids are also useful as competitive inhibitors of binding of the entire protein, or a portion thereof, to antibodies (as in

epitope mapping), and to natural binding partners, such as subunits in a multimeric complex or to receptors or ligands of the subject protein; this competitive inhibition permits identification and separation of molecules that bind specifically to the protein of interest, U.S. Pat. Nos. 5,539,084 and 5,783,674, incorporated herein by reference in their entireties.

The protein, or protein fragment, of the present invention is thus at least 6 amino acids in length, typically 10 at least 8, 9, 10 or 12 amino acids in length, and often at least 15 amino acids in length. Often, the protein or the present invention, or fragment thereof, is at least 20 amino acids in length, even 25 amino acids, 30 amino acids, 35 amino acids, or 50 amino acids or more in length. Of course, larger 15 fragments having at least 75 amino acids, 100 amino acids, or even 150 amino acids are also useful, and at times preferred.

The present invention further provides fusions of each of the proteins and protein fragments of the present invention to heterologous polypeptides.

By fusion is here intended that the protein or protein fragment of the present invention is linearly contiguous to the heterologous polypeptide in a peptide-bonded polymer of amino acids or amino acid analogues; by "heterologous polypeptide" is here intended a polypeptide that 25 does not naturally occur in contiguity with the protein or protein fragment of the present invention. As so defined, the fusion can consist entirely of a plurality of fragments of the RGL3 protein in altered arrangement; in such case, any of the RGL3 fragments can be considered heterologous to the other RGL3 30 fragments in the fusion protein. More typically, however, the heterologous polypeptide is not drawn from the RGL3 protein itself.

The fusion proteins of the present invention will include at least one fragment of the protein of the present invention, which fragment is at least 6, typically at least 8, often at least 15, and usefully at least 16, 17, 18, 19, or 20 amino acids long. The fragment of the protein of the present invention to be included in the fusion can usefully be at least 25 amino acids long, at least 50 amino acids long, and can be at least 75, 100, or even 150 amino acids long. Fusions that include the entirety of the proteins of the present invention have particular utility.

The heterologous polypeptide included within the fusion protein of the present invention is at least 6 amino acids in length, often at least 8 amino acids in length, and usefully at least 15, 20, and 25 amino acids in length. Fusions that include larger polypeptides, such as the IgG Fc region, and even entire proteins (such as GFP chromophore-containing proteins), have particular utility.

As described above in the description of vectors and expression vectors of the present invention, which discussion is incorporated herein by reference in its entirety, heterologous polypeptides to be included in the fusion proteins of the present invention can usefully include those designed to facilitate purification and/or visualization of recombinantly-expressed proteins. Although purification tags can also be incorporated into fusions that are chemically synthesized, chemical synthesis typically provides sufficient purity that further purification by HPLC suffices; however, visualization tags as above described retain their utility even when the protein is produced by chemical synthesis, and when so included render the fusion proteins of the present invention useful as directly detectable markers of RGL3 presence.

As also discussed above, heterologous polypeptides to be included in the fusion proteins of the present invention can

usefully include those that facilitate secretion of recombinantly expressed proteins – into the periplasmic space or extracellular milieu for prokaryotic hosts, into the culture medium for eukaryotic cells – through incorporation of 5 secretion signals and/or leader sequences.

Other useful protein fusions of the present invention include those that permit use of the protein of the present invention as bait in a yeast two-hybrid system. See Bartel et al. (eds.), The Yeast Two-Hybrid System, Oxford University Press (1997) (ISBN: 0195109384); Zhu et al., Yeast Hybrid Technologies, Eaton Publishing, (2000) (ISBN 1-881299-15-5); Fields et al., *Trends Genet.* 10(8):286-92 (1994); Mendelsohn et al., *Curr. Opin. Biotechnol.* 5(5):482-6 (1994); Luban et al., *Curr. Opin. Biotechnol.* 6(1):59-64 (1995); Allen et al., *Trends Biochem. Sci.* 20(12):511-6 (1995); Drees, *Curr. Opin. Chem. Biol.* 3(1):64-70 (1999); Topcu et al., *Pharm. Res.* 17(9):1049-55 (2000); Fashena et al., *Gene* 250(1-2):1-14 (2000), the disclosures of which are incorporated herein by reference in their entireties. Typically, such fusion is to either *E. coli* LexA or yeast GAL4 DNA binding domains. Related 20 bait plasmids are available that express the bait fused to a nuclear localization signal.

Other useful protein fusions include those that permit display of the encoded protein on the surface of a phage or cell, fusions to intrinsically fluorescent proteins, such as green fluorescent protein (GFP), and fusions to the IgG Fc region, as described above, which discussion is incorporated here by reference in its entirety.

The proteins and protein fragments of the present 30 invention can also usefully be fused to protein toxins, such as *Pseudomonas* exotoxin A, diphtheria toxin, shiga toxin A, anthrax toxin lethal factor, ricin, in order to effect ablation

of cells that bind or take up the proteins of the present invention.

The isolated proteins, protein fragments, and protein fusions of the present invention can be composed of natural 5 amino acids linked by native peptide bonds, or can contain any or all of nonnatural amino acid analogues, nonnative bonds, and post-synthetic (post translational) modifications, either throughout the length of the protein or localized to one or more portions thereof.

As is well known in the art, when the isolated protein is used, e.g., for epitope mapping, the range of such nonnatural analogues, nonnative inter-residue bonds, or post-synthesis modifications will be limited to those that permit binding of the peptide to antibodies. When used as an 10 immunogen for the preparation of antibodies in a non-human host, such as a mouse, the range of such nonnatural analogues, nonnative inter-residue bonds, or post-synthesis modifications will be limited to those that do not interfere with the 15 immunogenicity of the protein. When the isolated protein is used as a therapeutic agent, such as a vaccine or for replacement therapy, the range of such changes will be limited 20 to those that do not confer toxicity upon the isolated protein.

Non-natural amino acids can be incorporated during solid phase chemical synthesis or by recombinant techniques, 25 although the former is typically more common.

Solid phase chemical synthesis of peptides is well established in the art. Procedures are described, *inter alia*, in Chan et al. (eds.), Fmoc Solid Phase Peptide Synthesis: A Practical Approach (Practical Approach Series), Oxford Univ. Press (March 2000) (ISBN: 0199637245); Jones, Amino Acid and Peptide Synthesis (Oxford Chemistry Primers, No 7), Oxford Univ. Press (August 1992) (ISBN: 0198556683); and Bodanszky, Principles of Peptide Synthesis (Springer Laboratory), Springer

Verlag (December 1993) (ISBN: 0387564314), the disclosures of which are incorporated herein by reference in their entireties.

For example, D-enantiomers of natural amino acids can readily be incorporated during chemical peptide synthesis:

- 5 peptides assembled from D-amino acids are more resistant to proteolytic attack; incorporation of D-enantiomers can also be used to confer specific three dimensional conformations on the peptide. Other amino acid analogues commonly added during chemical synthesis include ornithine, norleucine,
- 10 phosphorylated amino acids (typically phosphoserine, phosphothreonine, phosphotyrosine), L-malonyltyrosine, a non-hydrolyzable analog of phosphotyrosine (Kole et al., *Biochem. Biophys. Res. Com.* 209:817-821 (1995)), and various halogenated phenylalanine derivatives.

15 Amino acid analogues having detectable labels are also usefully incorporated during synthesis to provide a labeled polypeptide.

Biotin, for example (indirectly detectable through interaction with avidin, streptavidin, neutravidin, captavidin, 20 or anti-biotin antibody), can be added using biotinoyl--(9-fluorenylmethoxycarbonyl)-L-lysine (FMOC biocytin) (Molecular Probes, Eugene, OR, USA). (Biotin can also be added enzymatically by incorporation into a fusion protein of a *E. coli* BirA substrate peptide.)

25 The FMOC and tBOC derivatives of dabcyl-L-lysine (Molecular Probes, Inc., Eugene, OR, USA) can be used to incorporate the dabcyl chromophore at selected sites in the peptide sequence during synthesis. The aminonaphthalene derivative EDANS, the most common fluorophore for pairing with 30 the dabcyl quencher in fluorescence resonance energy transfer (FRET) systems, can be introduced during automated synthesis of peptides by using EDANS--FMOC-L-glutamic acid or the corresponding tBOC derivative (both from Molecular Probes,

Inc., Eugene, OR, USA). Tetramethylrhodamine fluorophores can be incorporated during automated FMOC synthesis of peptides using (FMOC)--TMR-L-lysine (Molecular Probes, Inc. Eugene, OR, USA).

5 Other useful amino acid analogues that can be incorporated during chemical synthesis include aspartic acid, glutamic acid, lysine, and tyrosine analogues having allyl side-chain protection (Applied Biosystems, Inc., Foster City, CA, USA); the allyl side chain permits synthesis of cyclic, 10 branched-chain, sulfonated, glycosylated, and phosphorylated peptides.

A large number of other FMOC-protected non-natural amino acid analogues capable of incorporation during chemical synthesis are available commercially, including, e.g., Fmoc-2-aminobicyclo[2.2.1]heptane-2-carboxylic acid, Fmoc-3-endo-aminobicyclo[2.2.1]heptane-2-endo-carboxylic acid, Fmoc-3-exo-aminobicyclo[2.2.1]heptane-2-exo-carboxylic acid, Fmoc-3-endo-amino-bicyclo[2.2.1]hept-5-ene-2-endo-carboxylic acid, Fmoc-3-exo-amino-bicyclo[2.2.1]hept-5-ene-2-exo-carboxylic acid, Fmoc-20 cis-2-amino-1-cyclohexanecarboxylic acid, Fmoc-trans-2-amino-1-cyclohexanecarboxylic acid, Fmoc-1-amino-1-cyclopentanecarboxylic acid, Fmoc-cis-2-amino-1-cyclopentanecarboxylic acid, Fmoc-1-amino-1-cyclopropanecarboxylic acid, Fmoc-D-2-amino-4-(ethylthio)butyric acid, Fmoc-L-2-amino-4-(ethylthio)butyric acid, Fmoc-L-buthionine, Fmoc-S-methyl-L-Cysteine, Fmoc-2-aminobenzoic acid (anthranillic acid), Fmoc-3-aminobenzoic acid, Fmoc-4-aminobenzoic acid, Fmoc-2-aminobenzophenone-2'-carboxylic acid, Fmoc-N-(4-aminobenzoyl)-D-alanine, Fmoc-2-amino-4,5-dimethoxybenzoic acid, Fmoc-4-aminohippuric acid, Fmoc-2-amino-3-hydroxybenzoic acid, Fmoc-2-amino-5-hydroxybenzoic acid, Fmoc-3-amino-4-hydroxybenzoic acid, Fmoc-4-amino-3-hydroxybenzoic acid, Fmoc-4-amino-2-hydroxybenzoic

acid, Fmoc-5-amino-2-hydroxybenzoic acid, Fmoc-2-amino-3-methoxybenzoic acid, Fmoc-4-amino-3-methoxybenzoic acid, Fmoc-2-amino-3-methylbenzoic acid, Fmoc-2-amino-5-methylbenzoic acid, Fmoc-2-amino-6-methylbenzoic acid, Fmoc-3-amino-2-methylbenzoic acid, Fmoc-3-amino-4-methylbenzoic acid, Fmoc-4-amino-3-methylbenzoic acid, Fmoc-3-amino-2-naphtoic acid, Fmoc-D,L-3-amino-3-phenylpropionic acid, Fmoc-L-Methyldopa, Fmoc-2-amino-4,6-dimethyl-3-pyridinecarboxylic acid, Fmoc-D,L-?-amino-2-thiophenacetic acid, Fmoc-4-(carboxymethyl)piperazine, Fmoc-4-carboxypiperazine, Fmoc-4-(carboxymethyl)homopiperazine, Fmoc-4-phenyl-4-piperidinecarboxylic acid, Fmoc-L-1,2,3,4-tetrahydronorharman-3-carboxylic acid, Fmoc-L-thiazolidine-4-carboxylic acid, all available from The Peptide Laboratory (Richmond, CA, USA).

Non-natural residues can also be added biosynthetically by engineering a suppressor tRNA, typically one that recognizes the UAG stop codon, by chemical aminoacylation with the desired unnatural amino acid and. Conventional site-directed mutagenesis is used to introduce the chosen stop codon UAG at the site of interest in the protein gene. When the acylated suppressor tRNA and the mutant gene are combined in an *in vitro* transcription/translation system, the unnatural amino acid is incorporated in response to the UAG codon to give a protein containing that amino acid at the specified position. Liu et al., *Proc. Natl Acad. Sci. USA* 96(9):4780-5 (1999); Wang et al., *Science* 292(5516):498-500 (2001).

The isolated proteins, protein fragments and fusion proteins of the present invention can also include nonnative inter-residue bonds, including bonds that lead to circular and branched forms.

The isolated proteins and protein fragments of the present invention can also include post-translational and post-

synthetic modifications, either throughout the length of the protein or localized to one or more portions thereof.

For example, when produced by recombinant expression in eukaryotic cells, the isolated proteins, fragments, and fusion proteins of the present invention will typically include N-linked and/or O-linked glycosylation, the pattern of which will reflect both the availability of glycosylation sites on the protein sequence and the identity of the host cell. Further modification of glycosylation pattern can be performed enzymatically.

As another example, recombinant polypeptides of the invention may also include an initial modified methionine residue, in some cases resulting from host-mediated processes.

When the proteins, protein fragments, and protein fusions of the present invention are produced by chemical synthesis, post-synthetic modification can be performed before deprotection and cleavage from the resin or after deprotection and cleavage. Modification before deprotection and cleavage of the synthesized protein often allows greater control, e.g. by allowing targeting of the modifying moiety to the N-terminus of a resin-bound synthetic peptide.

Useful post-synthetic (and post-translational) modifications include conjugation to detectable labels, such as fluorophores.

A wide variety of amine-reactive and thiol-reactive fluorophore derivatives have been synthesized that react under nondenaturing conditions with N-terminal amino groups and epsilon amino groups of lysine residues, on the one hand, and with free thiol groups of cysteine residues, on the other.

Kits are available commercially that permit conjugation of proteins to a variety of amine-reactive or thiol-reactive fluorophores: Molecular Probes, Inc. (Eugene, OR, USA), e.g., offers kits for conjugating proteins to Alexa

Fluor 350, Alexa Fluor 430, Fluorescein-EX, Alexa Fluor 488, Oregon Green 488, Alexa Fluor 532, Alexa Fluor 546, Alexa Fluor 546, Alexa Fluor 568, Alexa Fluor 594, and Texas Red-X.

A wide variety of other amine-reactive and thiol-reactive fluorophores are available commercially (Molecular Probes, Inc., Eugene, OR, USA), including Alexa Fluor® 350, Alexa Fluor® 488, Alexa Fluor® 532, Alexa Fluor® 546, Alexa Fluor® 568, Alexa Fluor® 594, Alexa Fluor® 647 (monoclonal antibody labeling kits available from Molecular Probes, Inc., Eugene, OR, USA), BODIPY dyes, such as BODIPY 493/503, BODIPY FL, BODIPY R6G, BODIPY 530/550, BODIPY TMR, BODIPY 558/568, BODIPY 558/568, BODIPY 564/570, BODIPY 576/589, BODIPY 581/591, BODIPY TR, BODIPY 630/650, BODIPY 650/665, Cascade Blue, Cascade Yellow, Dansyl, lissamine rhodamine B, Marina Blue, Oregon Green 488, Oregon Green 514, Pacific Blue, rhodamine 6G, rhodamine green, rhodamine red, tetramethylrhodamine, Texas Red (available from Molecular Probes, Inc., Eugene, OR, USA).

The polypeptides of the present invention can also be conjugated to fluorophores, other proteins, and other macromolecules, using bifunctional linking reagents.

Common homobifunctional reagents include, e.g., APG, AEDP, BASED, BMB, BMDB, BMH, BMOE, BM[PEO]3, BM[PEO]4, BS3, BSOCOES, DFDNB, DMA, DMP, DMS, DPDPB, DSG, DSP (Lomant's Reagent), DSS, DST, DTBP, DTME, DTSSP, EGS, HBVS, 25 Sulfo-BSOCOES, Sulfo-DST, Sulfo-EGS (all available from Pierce, Rockford, IL, USA); common heterobifunctional cross-linkers include ABH, AMAS, ANB-NOS, APDP, ASBA, BMPA, BMPH, BMPS, EDC, EMCA, EMCH, EMCS, KMUA, KMUH, GMBS, LC-SMCC, LC-SPDP, MBS, M2C2H, MPBH, MSA, NHS-ASA, PDPH, PMPI, SADP, SAED, SAND, SANPAH, SASD, SATP, SBAP, SFAD, SIA, SIAB, SMCC, SMPB, SMPH, SMPT, SPDP, Sulfo-EMCS, Sulfo-GMBS, Sulfo-HSAB, Sulfo-KMUS, Sulfo-LC-SPDP, Sulfo-MBS, Sulfo-NHS-LC-ASA, Sulfo-SADP, Sulfo-SANPAH, Sulfo-SIAB, Sulfo-SMCC, Sulfo-SMPB,

Sulfo-LC-SMPT, SVSB, TFCS (all available Pierce, Rockford, IL, USA).

The proteins, protein fragments, and protein fusions of the present invention can be conjugated, using such cross-linking reagents, to fluorophores that are not amine- or thiol-reactive.

Other labels that usefully can be conjugated to the proteins, protein fragments, and fusion proteins of the present invention include radioactive labels, echosonographic contrast reagents, and MRI contrast agents.

The proteins, protein fragments, and protein fusions of the present invention can also usefully be conjugated using cross-linking agents to carrier proteins, such as KLH, bovine thyroglobulin, and even bovine serum albumin (BSA), to increase immunogenicity for raising anti-RGL3 antibodies.

The proteins, protein fragments, and protein fusions of the present invention can also usefully be conjugated to polyethylene glycol (PEG); PEGylation increases the serum half life of proteins administered intravenously for replacement therapy. Delgado *et al.*, *Crit. Rev. Ther. Drug Carrier Syst.* 9(3-4):249-304 (1992); Scott *et al.*, *Curr. Pharm. Des.* 4(6):423-38 (1998); De Santis *et al.*, *Curr. Opin. Biotechnol.* 10(4):324-30 (1999), incorporated herein by reference in their entireties. PEG monomers can be attached to the protein directly or through a linker, with PEGylation using PEG monomers activated with tresyl chloride (2,2,2-trifluoroethanesulphonyl chloride) permitting direct attachment under mild conditions.

The isolated proteins of the present invention, including fusions thereof, can be produced by recombinant expression, typically using the expression vectors of the present invention as above-described or, if fewer than about

100 amino acids, by chemical synthesis (typically, solid phase synthesis), and, on occasion, by *in vitro* translation.

Production of the isolated proteins of the present invention can optionally be followed by purification.

5 Purification of recombinantly expressed proteins is now well within the skill in the art. See, e.g., Thorner et al. (eds.), Applications of Chimeric Genes and Hybrid Proteins, Part A: Gene Expression and Protein Purification (Methods in Enzymology, Volume 326), Academic Press (2000), (ISBN: 10 0121822273); Harbin (ed.), Cloning, Gene Expression and Protein Purification : Experimental Procedures and Process Rationale, Oxford Univ. Press (2001) (ISBN: 0195132947); Marshak et al., Strategies for Protein Purification and Characterization: A Laboratory Course Manual, Cold Spring Harbor Laboratory Press 15 (1996) (ISBN: 0-87969-385-1); and Roe (ed.), Protein Purification Applications, Oxford University Press (2001), the disclosures of which are incorporated herein by reference in their entireties, and thus need not be detailed here.

Briefly, however, if purification tags have been 20 fused through use of an expression vector that appends such tag, purification can be effected, at least in part, by means appropriate to the tag, such as use of immobilized metal affinity chromatography for polyhistidine tags. Other techniques common in the art include ammonium sulfate 25 fractionation, immunoprecipitation, fast protein liquid chromatography (FPLC), high performance liquid chromatography (HPLC), and preparative gel electrophoresis.

Purification of chemically-synthesized peptides can readily be effected, e.g., by HPLC.

30 Accordingly, it is an aspect of the present invention to provide the isolated proteins of the present invention in pure or substantially pure form.

A purified protein of the present invention is an isolated protein, as above described, that is present at a concentration of at least 95%, as measured on a weight basis (w/w) with respect to total protein in a composition. Such purities can often be obtained during chemical synthesis without further purification, as, e.g., by HPLC. Purified proteins of the present invention can be present at a concentration (measured on a weight basis with respect to total protein in a composition) of 96%, 97%, 98%, and even 99%. The 10 proteins of the present invention can even be present at levels of 99.5%, 99.6%, and even 99.7%, 99.8%, or even 99.9% following purification, as by HPLC.

Although high levels of purity are particularly useful when the isolated proteins of the present invention are used as therapeutic agents — such as vaccines, or for replacement therapy — the isolated proteins of the present invention are also useful at lower purity. For example, partially purified proteins of the present invention can be used as immunogens to raise antibodies in laboratory animals.

Thus, in another aspect, the present invention provides the isolated proteins of the present invention in substantially purified form. A "substantially purified protein" of the present invention is an isolated protein, as above described, present at a concentration of at least 70%, measured on a weight basis with respect to total protein in a composition. Usefully, the substantially purified protein is present at a concentration, measured on a weight basis with respect to total protein in a composition, of at least 75%, 80%, or even at least 85%, 90%, 91%, 92%, 93%, 94%, 94.5% or 30 even at least 94.9%.

In preferred embodiments, the purified and substantially purified proteins of the present invention are in

compositions that lack detectable amphotytes, acrylamide monomers, bis-acrylamide monomers, and polyacrylamide.

The proteins, fragments, and fusions of the present invention can usefully be attached to a substrate. The 5 substrate can porous or solid, planar or non-planar; the bond can be covalent or noncovalent.

For example, the proteins, fragments, and fusions of the present invention can usefully be bound to a porous substrate, commonly a membrane, typically comprising 10 nitrocellulose, polyvinylidene fluoride (PVDF), or cationically derivatized, hydrophilic PVDF; so bound, the proteins, fragments, and fusions of the present invention can be used to detect and quantify antibodies, e.g. in serum, that bind specifically to the immobilized protein of the present 15 invention.

As another example, the proteins, fragments, and fusions of the present invention can usefully be bound to a substantially nonporous substrate, such as plastic, to detect and quantify antibodies, e.g. in serum, that bind specifically 20 to the immobilized protein of the present invention. Such plastics include polymethylacrylic, polyethylene, polypropylene, polyacrylate, polymethylmethacrylate, polyvinylchloride, polytetrafluoroethylene, polystyrene, polycarbonate, polyacetal, polysulfone, celluloseacetate, 25 cellulosenitrate, nitrocellulose, or mixtures thereof; when the assay is performed in standard microtiter dish, the plastic is typically polystyrene.

The proteins, fragments, and fusions of the present invention can also be attached to a substrate suitable for use 30 as a surface enhanced laser desorption ionization source; so attached, the protein, fragment, or fusion of the present invention is useful for binding and then detecting secondary proteins that bind with sufficient affinity or avidity to the

surface-bound protein to indicate biologic interaction therebetween. The proteins, fragments, and fusions of the present invention can also be attached to a substrate suitable for use in surface plasmon resonance detection; so attached, 5 the protein, fragment, or fusion of the present invention is useful for binding and then detecting secondary proteins that bind with sufficient affinity or avidity to the surface-bound protein to indicate biological interaction therebetween.

10 RGL3 Proteins

In a first series of protein embodiments, the invention provides an isolated RGL3 polypeptide having the amino acid sequence in SEQ ID NO: 3, which is full length RGL3 15 proteins. When used as immunogens, the full length proteins of the present invention can be used, *inter alia*, to elicit antibodies that bind to a variety of epitopes of the RGL3 protein.

The invention further provides fragments of the 20 above-described polypeptides, particularly fragments having at least 6 amino acids, typically at least 8 amino acids, often at least 15 amino acids, and even the entirety of the sequence given in SEQ ID NO: 3.

As described above, the invention further provides 25 proteins that differ in sequence from those described with particularity in the above-referenced SEQ ID NOs., whether by way of insertion or deletion, by way of conservative or moderately conservative substitutions, as hybridization related proteins, or as cross-hybridizing proteins, with those that 30 substantially retain a RGL3 activity particularly useful.

The invention further provides fusions of the proteins and protein fragments herein described to heterologous polypeptides.

ANTIBODIES AND ANTIBODY-PRODUCING CELLS

In another aspect, the invention provides antibodies,
5 including fragments and derivatives thereof, that bind
specifically to RGL3 proteins and protein fragments of the
present invention or to one or more of the proteins and protein
fragments encoded by the isolated RGL3 nucleic acids of the
present invention. The antibodies of the present invention can
10 be specific for all of linear epitopes, discontinuous epitopes,
or conformational epitopes of such proteins or protein
fragments, either as present on the protein in its native
conformation or, in some cases, as present on the proteins as
denatured, as, e.g., by solubilization in SDS.

15 In other embodiments, the invention provides
antibodies, including fragments and derivatives thereof, the
binding of which can be competitively inhibited by one or more
of the RGL3 proteins and protein fragments of the present
invention, or by one or more of the proteins and protein
20 fragments encoded by the isolated RGL3 nucleic acids of the
present invention.

As used herein, the term "antibody" refers to a
polypeptide, at least a portion of which is encoded by at least
one immunoglobulin gene, which can bind specifically to a first
25 molecular species, and to fragments or derivatives thereof that
remain capable of such specific binding.

By "bind specifically" and "specific binding" is here
intended the ability of the antibody to bind to a first
molecular species in preference to binding to other molecular
30 species with which the antibody and first molecular species are
admixed. An antibody is said specifically to "recognize" a
first molecular species when it can bind specifically to that
first molecular species.

As is well known in the art, the degree to which an antibody can discriminate as among molecular species in a mixture will depend, in part, upon the conformational relatedness of the species in the mixture; typically, the 5 antibodies of the present invention will discriminate over adventitious binding to non-RGL3 proteins by at least two-fold, more typically by at least 5-fold, typically by more than 10-fold, 25-fold, 50-fold, 75-fold, and often by more than 100-fold, and on occasion by more than 500-fold or 1000-fold. When 10 used to detect the proteins or protein fragments of the present invention, the antibody of the present invention is sufficiently specific when it can be used to determine the presence of the protein of the present invention in samples derived from human adrenal, adult liver, bone marrow, brain, 15 fetal liver, heart, kidney, lung, placenta, colon, skeletal muscle and prostate.

Typically, the affinity or avidity of an antibody (or antibody multimer, as in the case of an IgM pentamer) of the present invention for a protein or protein fragment of the 20 present invention will be at least about 1×10^{-6} molar (M), typically at least about 5×10^{-7} M, usefully at least about 1×10^{-7} M, with affinities and avidities of at least 1×10^{-8} M, 5×10^{-9} M, and 1×10^{-10} M proving especially useful.

The antibodies of the present invention can be 25 naturally-occurring forms, such as IgG, IgM, IgD, IgE, and IgA, from any mammalian species.

Human antibodies can, but will infrequently, be drawn directly from human donors or human cells. In such case, antibodies to the proteins of the present invention will 30 typically have resulted from fortuitous immunization, such as autoimmune immunization, with the protein or protein fragments of the present invention. Such antibodies will typically, but will not invariably, be polyclonal.

Human antibodies are more frequently obtained using transgenic animals that express human immunoglobulin genes, which transgenic animals can be affirmatively immunized with the protein immunogen of the present invention. Human Ig-
5 transgenic mice capable of producing human antibodies and methods of producing human antibodies therefrom upon specific immunization are described, *inter alia*, in U.S. Patent Nos. 6,162,963; 6,150,584; 6,114,598; 6,075,181; 5,939,598; 5,877,397; 5,874,299; 5,814,318; 5,789,650; 5,770,429;
10 5,661,016; 5,633,425; 5,625,126; 5,569,825; 5,545,807; 5,545,806, and 5,591,669, the disclosures of which are incorporated herein by reference in their entireties. Such antibodies are typically monoclonal, and are typically produced using techniques developed for production of murine antibodies.
15 Human antibodies are particularly useful, and often preferred, when the antibodies of the present invention are to be administered to human beings as *in vivo* diagnostic or therapeutic agents, since recipient immune response to the administered antibody will often be substantially less than
20 that occasioned by administration of an antibody derived from another species, such as mouse.

IgG, IgM, IgD, IgE and IgA antibodies of the present invention are also usefully obtained from other mammalian species, including rodents - typically mouse, but also rat,
25 guinea pig, and hamster - lagomorphs, typically rabbits, and also larger mammals, such as sheep, goats, cows, and horses. In such cases, as with the transgenic human-antibody-producing non-human mammals, fortuitous immunization is not required, and the non-human mammal is typically affirmatively immunized,
30 according to standard immunization protocols, with the protein or protein fragment of the present invention.

As discussed above, virtually all fragments of 8 or more contiguous amino acids of the proteins of the present

invention can be used effectively as immunogens when conjugated to a carrier, typically a protein such as bovine thyroglobulin, keyhole limpet hemocyanin, or bovine serum albumin, conveniently using a bifunctional linker such as those
5 described elsewhere above, which discussion is incorporated by reference here.

Immunogenicity can also be conferred by fusion of the proteins and protein fragments of the present invention to other moieties.

10 For example, peptides of the present invention can be produced by solid phase synthesis on a branched polylysine core matrix; these multiple antigenic peptides (MAPs) provide high purity, increased avidity, accurate chemical definition and improved safety in vaccine development. Tam et al., Proc.
15 Natl. Acad. Sci. USA 85:5409-5413 (1988); Posnett et al., J. Biol. Chem. 263, 1719-1725 (1988).

Protocols for immunizing non-human mammals are well-established in the art, Harlow et al. (eds.), Antibodies: A Laboratory Manual, Cold Spring Harbor Laboratory (1998) (ISBN: 20 0879693142); Coligan et al. (eds.), Current Protocols in Immunology, John Wiley & Sons, Inc. (2001) (ISBN: 0-471-52276-7); Zola, Monoclonal Antibodies : Preparation and Use of Monoclonal Antibodies and Engineered Antibody Derivatives (Basics: From Background to Bench), Springer Verlag 25 (2000) (ISBN: 0387915907), the disclosures of which are incorporated herein by reference, and often include multiple immunizations, either with or without adjuvants such as Freund's complete adjuvant and Freund's incomplete adjuvant.

Antibodies from nonhuman mammals can be polyclonal or 30 monoclonal, with polyclonal antibodies having certain advantages in immunohistochemical detection of the proteins of the present invention and monoclonal antibodies having

advantages in identifying and distinguishing particular epitopes of the proteins of the present invention.

Following immunization, the antibodies of the present invention can be produced using any art-accepted technique.

- 5 Such techniques are well known in the art, Coligan *et al.* (eds.), Current Protocols in Immunology, John Wiley & Sons, Inc. (2001) (ISBN: 0-471-52276-7); Zola, Monoclonal Antibodies: Preparation and Use of Monoclonal Antibodies and Engineered Antibody Derivatives (Basics: From Background to Bench),
- 10 Springer Verlag (2000) (ISBN: 0387915907); Howard *et al.* (eds.), Basic Methods in Antibody Production and Characterization, CRC Press (2000) (ISBN: 0849394457); Harlow *et al.* (eds.), Antibodies: A Laboratory Manual, Cold Spring Harbor Laboratory (1998) (ISBN: 0879693142); Davis (ed.),
- 15 Monoclonal Antibody Protocols, Vol. 45, Humana Press (1995) (ISBN: 0896033082); Delves (ed.), Antibody Production: Essential Techniques, John Wiley & Son Ltd (1997) (ISBN: 0471970107); Kenney, Antibody Solution: An Antibody Methods Manual, Chapman & Hall (1997) (ISBN: 0412141914), incorporated
20 herein by reference in their entireties, and thus need not be detailed here.

Briefly, however, such techniques include, *inter alia*, production of monoclonal antibodies by hybridomas and expression of antibodies or fragments or derivatives thereof
25 from host cells engineered to express immunoglobulin genes or fragments thereof. These two methods of production are not mutually exclusive: genes encoding antibodies specific for the proteins or protein fragments of the present invention can be cloned from hybridomas and thereafter expressed in other host
30 cells. Nor need the two necessarily be performed together: e.g., genes encoding antibodies specific for the proteins and protein fragments of the present invention can be cloned directly from B cells known to be specific for the desired

protein, as further described in U.S. Pat. No. 5,627,052, the disclosure of which is incorporated herein by reference in its entirety, or from antibody-displaying phage.

Recombinant expression in host cells is particularly useful when fragments or derivatives of the antibodies of the present invention are desired.

Host cells for recombinant antibody production – either whole antibodies, antibody fragments, or antibody derivatives – can be prokaryotic or eukaryotic.

Prokaryotic hosts are particularly useful for producing phage displayed antibodies of the present invention.

The technology of phage-displayed antibodies, in which antibody variable region fragments are fused, for example, to the gene III protein (pIII) or gene VIII protein (pVIII) for display on the surface of filamentous phage, such as M13, is by now well-established, Sidhu, *Curr. Opin. Biotechnol.* 11(6):610-6 (2000); Griffiths et al., *Curr. Opin. Biotechnol.* 9(1):102-8 (1998); Hoogenboom et al., *Current Immunotechnology*, 4(1):1-20 (1998); Rader et al., *Current Opinion in Biotechnology* 8:503-508 (1997); Aujame et al., *Human Antibodies* 8:155-168 (1997); Hoogenboom, *Trends in Biotechnol.* 15:62-70 (1997); de Kruif et al., 17:453-455 (1996); Barbas et al., *Trends in Biotechnol.* 14:230-234 (1996); Winter et al., *Ann. Rev. Immunol.* 433-455 (1994), and techniques and protocols required to generate, propagate, screen (pan), and use the antibody fragments from such libraries have recently been compiled, Barbas et al., Phage Display: A Laboratory Manual, Cold Spring Harbor Laboratory Press (2001) (ISBN 0-87969-546-3); Kay et al. (eds.), Phage Display of Peptides and Proteins: A Laboratory Manual, Academic Press, Inc. (1996); Abelson et al. (eds.), Combinatorial Chemistry, Methods in Enzymology vol. 267, Academic Press (May 1996), the disclosures

of which are incorporated herein by reference in their entireties.

Typically, phage-displayed antibody fragments are scFv fragments or Fab fragments; when desired, full length antibodies can be produced by cloning the variable regions from the displaying phage into a complete antibody and expressing the full length antibody in a further prokaryotic or a eukaryotic host cell.

Eukaryotic cells are also useful for expression of the antibodies, antibody fragments, and antibody derivatives of the present invention.

For example, antibody fragments of the present invention can be produced in *Pichia pastoris*, Takahashi et al., *Biosci. Biotechnol. Biochem.* 64(10):2138-44 (2000); Freyre et al., *J. Biotechnol.* 76(2-3):157-63 (2000); Fischer et al., *Biotechnol. Appl. Biochem.* 30 (Pt 2):117-20 (1999); Pennell et al., *Res. Immunol.* 149(6):599-603 (1998); Eldin et al., *J. Immunol. Methods.* 201(1):67-75 (1997); and in *Saccharomyces cerevisiae*, Frenken et al., *Res. Immunol.* 149(6):589-99 (1998); Shusta et al., *Nature Biotechnol.* 16(8):773-7 (1998), the disclosures of which are incorporated herein by reference in their entireties.

Antibodies, including antibody fragments and derivatives, of the present invention can also be produced in insect cells, Li et al., *Protein Expr. Purif.* 21(1):121-8 (2001); Ailor et al., *Biotechnol. Bioeng.* 58(2-3):196-203 (1998); Hsu et al., *Biotechnol. Prog.* 13(1):96-104 (1997); Edelman et al., *Immunology* 91(1):13-9 (1997); and Nesbit et al., *J. Immunol. Methods.* 151(1-2):201-8 (1992), the disclosures of which are incorporated herein by reference in their entireties.

Antibodies and fragments and derivatives thereof of the present invention can also be produced in plant cells,

Giddings et al., *Nature Biotechnol.* 18(11):1151-5 (2000); Gavilondo et al., *Biotechniques* 29(1):128-38 (2000); Fischer et al., *J. Biol. Regul. Homeost. Agents* 14(2):83-92 (2000); Fischer et al., *Biotechnol. Appl. Biochem.* 30 (Pt 2):113-6 5 (1999); Fischer et al., *Biol. Chem.* 380(7-8):825-39 (1999); Russell, *Curr. Top. Microbiol. Immunol.* 240:119-38 (1999); and Ma et al., *Plant Physiol.* 109(2):341-6 (1995), the disclosures of which are incorporated herein by reference in their entireties.

10 Mammalian cells useful for recombinant expression of antibodies, antibody fragments, and antibody derivatives of the present invention include CHO cells, COS cells, 293 cells, and myeloma cells.

15 Verma et al., *J. Immunol. Methods* 216(1-2):165-81 (1998), review and compare bacterial, yeast, insect and mammalian expression systems for expression of antibodies.

20 Antibodies of the present invention can also be prepared by cell free translation, as further described in Merk et al., *J. Biochem.* (Tokyo). 125(2):328-33 (1999) and Ryabova et al., *Nature Biotechnol.* 15(1):79-84 (1997), and in the milk of transgenic animals, as further described in Pollock et al., *J. Immunol. Methods* 231(1-2):147-57 (1999), the disclosures of which are incorporated herein by reference in their entireties.

25 The invention further provides antibody fragments that bind specifically to one or more of the proteins and protein fragments of the present invention, to one or more of the proteins and protein fragments encoded by the isolated nucleic acids of the present invention, or the binding of which can be competitively inhibited by one or more of the proteins 30 and protein fragments of the present invention or one or more of the proteins and protein fragments encoded by the isolated nucleic acids of the present invention.

Among such useful fragments are Fab, Fab', Fv, F(ab)'₂, and single chain Fv (scFv) fragments. Other useful fragments are described in Hudson, *Curr. Opin. Biotechnol.* 9(4):395-402 (1998).

5 It is also an aspect of the present invention to provide antibody derivatives that bind specifically to one or more of the proteins and protein fragments of the present invention, to one or more of the proteins and protein fragments encoded by the isolated nucleic acids of the present invention,
10 or the binding of which can be competitively inhibited by one or more of the proteins and protein fragments of the present invention or one or more of the proteins and protein fragments encoded by the isolated nucleic acids of the present invention.

15 Among such useful derivatives are chimeric, primateized, and humanized antibodies; such derivatives are less immunogenic in human beings, and thus more suitable for in vivo administration, than are unmodified antibodies from non-human mammalian species.

20 Chimeric antibodies typically include heavy and/or light chain variable regions (including both CDR and framework residues) of immunoglobulins of one species, typically mouse, fused to constant regions of another species, typically human.

25 See, e.g., U.S. Pat. No. 5,807,715; Morrison et al., *Proc. Natl. Acad. Sci USA* 81(21):6851-5 (1984); Sharon et al., *Nature* 309(5966):364-7 (1984); Takeda et al., *Nature* 314(6010):452-4 (1985), the disclosures of which are incorporated herein by reference in their entireties. Primateized and humanized antibodies typically include heavy and/or light chain CDRs from a murine antibody grafted into a non-human primate or human
30 antibody V region framework, usually further comprising a human constant region, Riechmann et al., *Nature* 332(6162):323-7 (1988); Co et al., *Nature* 351(6326):501-2 (1991); U.S. Pat. Nos. 6,054,297; 5,821,337; 5,770,196; 5,766,886; 5,821,123;

5,869,619; 6,180,377; 6,013,256; 5,693,761; and 6,180,370, the disclosures of which are incorporated herein by reference in their entireties.

Other useful antibody derivatives of the invention 5 include heteromeric antibody complexes and antibody fusions, such as diabodies (bispecific antibodies), single-chain diabodies, and intrabodies.

The antibodies of the present invention, including fragments and derivatives thereof, can usefully be labeled. It 10 is, therefore, another aspect of the present invention to provide labeled antibodies that bind specifically to one or more of the proteins and protein fragments of the present invention, to one or more of the proteins and protein fragments encoded by the isolated nucleic acids of the present invention, 15 or the binding of which can be competitively inhibited by one or more of the proteins and protein fragments of the present invention or one or more of the proteins and protein fragments encoded by the isolated nucleic acids of the present invention.

The choice of label depends, in part, upon the 20 desired use.

For example, when the antibodies of the present invention are used for immunohistochemical staining of tissue samples, the label can usefully be an enzyme that catalyzes production and local deposition of a detectable product.

Enzymes typically conjugated to antibodies to permit 25 their immunohistochemical visualization are well known, and include alkaline phosphatase, β -galactosidase, glucose oxidase, horseradish peroxidase (HRP), and urease. Typical substrates for production and deposition of visually detectable products 30 include o-nitrophenyl-beta-D-galactopyranoside (ONPG); o-phenylenediamine dihydrochloride (OPD); p-nitrophenyl phosphate (PNPP); p-nitrophenyl-beta-D-galactopyranoside (PNPG); 3',3'-diaminobenzidine (DAB); 3-amino-9-ethylcarbazole

(AEC) ; 4-chloro-1-naphthol (CN) ;
5-bromo-4-chloro-3-indolyl-phosphate (BCIP) ; ABTS®; Bluogal ;
iodonitrotetrazolium (INT) ; nitroblue tetrazolium chloride
(NBT) ; phenazine methosulfate (PMS) ; phenolphthalein
5 monophosphate (PMP) ; tetramethyl benzidine (TMB) ;
tetranitroblue tetrazolium (TNBT) ; X-Gal ; X-Gluc ; and
X-Glucoside.

Other substrates can be used to produce products for local deposition that are luminescent. For example, in the
10 presence of hydrogen peroxide (H_2O_2) , horseradish peroxidase (HRP) can catalyze the oxidation of cyclic diacylhydrazides, such as luminol. Immediately following the oxidation, the luminol is in an excited state (intermediate reaction product), which decays to the ground state by emitting light. Strong
15 enhancement of the light emission is produced by enhancers, such as phenolic compounds. Advantages include high sensitivity, high resolution, and rapid detection without radioactivity and requiring only small amounts of antibody.
See, e.g., Thorpe et al., *Methods Enzymol.* 133:331-53 (1986) ;
20 Kricka et al., *J. Immunoassay* 17(1):67-83 (1996) ; and Lundqvist et al., *J. Biolumin. Chemilumin.* 10(6):353-9 (1995) , the disclosures of which are incorporated herein by reference in their entireties. Kits for such enhanced chemiluminescent detection (ECL) are available commercially.

25 The antibodies can also be labeled using colloidal gold.

As another example, when the antibodies of the present invention are used, e.g., for flow cytometric detection, for scanning laser cytometric detection, or for
30 fluorescent immunoassay, they can usefully be labeled with fluorophores.

There are a wide variety of fluorophore labels that can usefully be attached to the antibodies of the present invention.

For flow cytometric applications, both for extracellular detection and for intracellular detection, common useful fluorophores can be fluorescein isothiocyanate (FITC), allophycocyanin (APC), R-phycoerythrin (PE), peridinin chlorophyll protein (PerCP), Texas Red, Cy3, Cy5, fluorescence resonance energy tandem fluorophores such as PerCP-Cy5.5, PE-Cy5, PE-Cy5.5, PE-Texas Red, and APC-Cy7.

Other fluorophores include, *inter alia*, Alexa Fluor® 350, Alexa Fluor® 488, Alexa Fluor® 532, Alexa Fluor® 546, Alexa Fluor® 568, Alexa Fluor® 594, Alexa Fluor® 647 (monoclonal antibody labeling kits available from Molecular Probes, Inc., Eugene, OR, USA), BODIPY dyes, such as BODIPY 493/503, BODIPY FL, BODIPY R6G, BODIPY 530/550, BODIPY TMR, BODIPY 558/568, BODIPY 558/568, BODIPY 564/570, BODIPY 576/589, BODIPY 581/591, BODIPY TR, BODIPY 630/650, BODIPY 650/665, Cascade Blue, Cascade Yellow, Dansyl, lissamine rhodamine B, Marina Blue, Oregon Green 488, Oregon Green 514, Pacific Blue, rhodamine 6G, rhodamine green, rhodamine red, tetramethylrhodamine, Texas Red (available from Molecular Probes, Inc., Eugene, OR, USA), and Cy2, Cy3, Cy3.5, Cy5, Cy5.5, Cy7, all of which are also useful for fluorescently labeling the antibodies of the present invention.

For secondary detection using labeled avidin, streptavidin, captavidin or neutravidin, the antibodies of the present invention can usefully be labeled with biotin.

When the antibodies of the present invention are used, e.g., for western blotting applications, they can usefully be labeled with radioisotopes, such as ^{33}P , ^{32}P , ^{35}S , ^3H , and ^{125}I .

As another example, when the antibodies of the present invention are used for radioimmunotherapy, the label can usefully be ^{228}Th , ^{227}Ac , ^{225}Ac , ^{223}Ra , ^{213}Bi , ^{212}Pb , ^{212}Bi , ^{211}At , ^{203}Pb , ^{194}Os , ^{188}Re , ^{186}Re , ^{153}Sm , ^{149}Tb , ^{131}I , ^{125}I , ^{111}In , ^{105}Rh , ^{99m}Tc , 5 ^{97}Ru , ^{90}Y , ^{90}Sr , ^{88}Y , ^{72}Se , ^{67}Cu , or ^{47}Sc .

As another example, when the antibodies of the present invention are to be used for *in vivo* diagnostic use, they can be rendered detectable by conjugation to MRI contrast agents, such as gadolinium diethylenetriaminepentaacetic acid (DTPA), Lauffer et al., *Radiology* 207(2):529-38 (1998), or by 10 radioisotopic labeling.

As would be understood, use of the labels described above is not restricted to the application as for which they were mentioned.

The antibodies of the present invention, including 15 fragments and derivatives thereof, can also be conjugated to toxins, in order to target the toxin's ablative action to cells that display and/or express the proteins of the present invention. Commonly, the antibody in such immunotoxins is conjugated to *Pseudomonas exotoxin A*, diphtheria toxin, shiga 20 toxin A, anthrax toxin lethal factor, or ricin. See Hall (ed.), Immunotoxin Methods and Protocols (Methods in Molecular Biology, Vol 166), Humana Press (2000) (ISBN:0896037754); and Frankel et al. (eds.), Clinical Applications of Immunotoxins, 25 Springer-Verlag New York, Incorporated (1998) (ISBN:3540640975), the disclosures of which are incorporated herein by reference in their entireties, for review.

The antibodies of the present invention can usefully be attached to a substrate, and it is, therefore, another 30 aspect of the invention to provide antibodies that bind specifically to one or more of the proteins and protein fragments of the present invention, to one or more of the proteins and protein fragments encoded by the isolated nucleic

acids of the present invention, or the binding of which can be competitively inhibited by one or more of the proteins and protein fragments of the present invention or one or more of the proteins and protein fragments encoded by the isolated 5 nucleic acids of the present invention, attached to a substrate.

Substrates can be porous or nonporous, planar or nonplanar.

For example, the antibodies of the present invention 10 can usefully be conjugated to filtration media, such as NHS-activated Sepharose or CNBr-activated Sepharose for purposes of immunoaffinity chromatography.

For example, the antibodies of the present invention can usefully be attached to paramagnetic microspheres, 15 typically by biotin-streptavidin interaction, which microsphere can then be used for isolation of cells that express or display the proteins of the present invention. As another example, the antibodies of the present invention can usefully be attached to the surface of a microtiter plate for ELISA.

As noted above, the antibodies of the present 20 invention can be produced in prokaryotic and eukaryotic cells.

It is, therefore, another aspect of the present invention to provide cells that express the antibodies of the present invention, including hybridoma cells, B cells, plasma cells, 25 and host cells recombinantly modified to express the antibodies of the present invention.

In yet a further aspect, the present invention provides aptamers evolved to bind specifically to one or more of the proteins and protein fragments of the present invention, 30 to one or more of the proteins and protein fragments encoded by the isolated nucleic acids of the present invention, or the binding of which can be competitively inhibited by one or more of the proteins and protein fragments of the present invention

or one or more of the proteins and protein fragments encoded by the isolated nucleic acids of the present invention.

RGL3 Antibodies

5

In a first series of antibody embodiments, the invention provides antibodies, both polyclonal and monoclonal, and fragments and derivatives thereof, that bind specifically to a polypeptide having the amino acid sequence in SEQ ID NO: 10 3, which is full length RGL3 protein.

Such antibodies are useful in *in vitro* immunoassays, such as ELISA, western blot or immunohistochemical assay of tumor tissue or cells. Such antibodies are also useful in isolating and purifying RGL3 proteins, including related cross-reactive proteins, by immunoprecipitation, immunoaffinity chromatography, or magnetic bead-mediated purification. 15

In another series of antibody embodiments, the invention provides antibodies, both polyclonal and monoclonal, and fragments and derivatives thereof, the specific binding of 20 which can be competitively inhibited by the isolated proteins and polypeptides of the present invention.

In other embodiments, the invention further provides the above-described antibodies detectably labeled, and in yet other embodiments, provides the above-described antibodies 25 attached to a substrate.

PHARMACEUTICAL COMPOSITIONS

RGL3 is important as a guanine nucleotide exchange 30 factor for the small GTPase Ral and a downstream effector for both Rit and Ras. Defects in RGL3 expression, activity, distribution, localization, and/or solubility are a cause of human disease, which disease can manifest as a disorder of

adrenal, adult liver, bone marrow, brain, fetal liver, heart, kidney, lung, placenta, colon, skeletal muscle and prostate function.

Accordingly, pharmaceutical compositions comprising
5 nucleic acids, proteins, and antibodies of the present invention, as well as mimetics, agonists, antagonists, or inhibitors of RGL3 activity, can be administered as therapeutics for treatment of RGL3 defects.

Thus, in another aspect, the invention provides
10 pharmaceutical compositions comprising the nucleic acids, nucleic acid fragments, proteins, protein fusions, protein fragments, antibodies, antibody derivatives, antibody fragments, mimetics, agonists, antagonists, and inhibitors of the present invention.

15 Such a composition typically contains from about 0.1 to 90% by weight of a therapeutic agent of the invention formulated in and/or with a pharmaceutically acceptable carrier or excipient.

Pharmaceutical formulation is a well-established art,
20 and is further described in Gennaro (ed.), Remington: The Science and Practice of Pharmacy, 20th ed., Lippincott, Williams & Wilkins (2000) (ISBN: 0683306472); Ansel et al., Pharmaceutical Dosage Forms and Drug Delivery Systems, 7th ed., Lippincott Williams & Wilkins Publishers (1999) (ISBN:
25 0683305727); and Kibbe (ed.), Handbook of Pharmaceutical Excipients American Pharmaceutical Association, 3rd ed. (2000) (ISBN: 091733096X), the disclosures of which are incorporated herein by reference in their entireties, and thus need not be described in detail herein.

30 Briefly, however, formulation of the pharmaceutical compositions of the present invention will depend upon the route chosen for administration. The pharmaceutical compositions utilized in this invention can be administered by

various routes including both enteral and parenteral routes, including oral, intravenous, intramuscular, subcutaneous, inhalation, topical, sublingual, rectal, intra-arterial, intramedullary, intrathecal, intraventricular, transmucosal, 5 transdermal, intranasal, intraperitoneal, intrapulmonary, and intrauterine.

Oral dosage forms can be formulated as tablets, pills, dragees, capsules, liquids, gels, syrups, slurries, suspensions, and the like, for ingestion by the patient.

10 Solid formulations of the compositions for oral administration can contain suitable carriers or excipients, such as carbohydrate or protein fillers, such as sugars, including lactose, sucrose, mannitol, or sorbitol; starch from corn, wheat, rice, potato, or other plants; cellulose, such as 15 methyl cellulose, hydroxypropylmethyl-cellulose, sodium carboxymethylcellulose, or microcrystalline cellulose; gums including arabic and tragacanth; proteins such as gelatin and collagen; inorganics, such as kaolin, calcium carbonate, dicalcium phosphate, sodium chloride; and other agents such as 20 acacia and alginic acid.

Agents that facilitate disintegration and/or solubilization can be added, such as the cross-linked polyvinyl pyrrolidone, agar, alginic acid, or a salt thereof, such as sodium alginate, microcrystalline cellulose, corn starch, 25 sodium starch glycolate, and alginic acid.

Tablet binders that can be used include acacia, methylcellulose, sodium carboxymethylcellulose, polyvinylpyrrolidone (PovidoneTM), hydroxypropyl methylcellulose, sucrose, starch and ethylcellulose.

30 Lubricants that can be used include magnesium stearates, stearic acid, silicone fluid, talc, waxes, oils, and colloidal silica.

Fillers, agents that facilitate disintegration and/or solubilization, tablet binders and lubricants, including the aforementioned, can be used singly or in combination.

Solid oral dosage forms need not be uniform
5 throughout.

For example, dragee cores can be used in conjunction with suitable coatings, such as concentrated sugar solutions, which can also contain gum arabic, talc, polyvinylpyrrolidone, carbopol gel, polyethylene glycol, and/or titanium dioxide, 10 lacquer solutions, and suitable organic solvents or solvent mixtures.

Oral dosage forms of the present invention include push-fit capsules made of gelatin, as well as soft, sealed capsules made of gelatin and a coating, such as glycerol or 15 sorbitol. Push-fit capsules can contain active ingredients mixed with a filler or binders, such as lactose or starches, lubricants, such as talc or magnesium stearate, and, optionally, stabilizers. In soft capsules, the active compounds can be dissolved or suspended in suitable liquids, 20 such as fatty oils, liquid, or liquid polyethylene glycol with or without stabilizers.

Additionally, dyestuffs or pigments can be added to the tablets or dragee coatings for product identification or to characterize the quantity of active compound, i.e., dosage.

Liquid formulations of the pharmaceutical 25 compositions for oral (enteral) administration are prepared in water or other aqueous vehicles and can contain various suspending agents such as methylcellulose, alginates, tragacanth, pectin, kelgin, carrageenan, acacia, 30 polyvinylpyrrolidone, and polyvinyl alcohol. The liquid formulations can also include solutions, emulsions, syrups and elixirs containing, together with the active compound(s), wetting agents, sweeteners, and coloring and flavoring agents.

The pharmaceutical compositions of the present invention can also be formulated for parenteral administration.

For intravenous injection, water soluble versions of the compounds of the present invention are formulated in, or if provided as a lyophilate, mixed with, a physiologically acceptable fluid vehicle, such as 5% dextrose ("D5"), physiologically buffered saline, 0.9% saline, Hanks' solution, or Ringer's solution.

Intramuscular preparations, e.g. a sterile formulation of a suitable soluble salt form of the compounds of the present invention, can be dissolved and administered in a pharmaceutical excipient such as Water-for-Injection, 0.9% saline, or 5% glucose solution. Alternatively, a suitable insoluble form of the compound can be prepared and administered as a suspension in an aqueous base or a pharmaceutically acceptable oil base, such as an ester of a long chain fatty acid (e.g., ethyl oleate), fatty oils such as sesame oil, triglycerides, or liposomes.

Parenteral formulations of the compositions can contain various carriers such as vegetable oils, dimethylacetamide, dimethylformamide, ethyl lactate, ethyl carbonate, isopropyl myristate, ethanol, polyols (glycerol, propylene glycol, liquid polyethylene glycol, and the like).

Aqueous injection suspensions can also contain substances that increase the viscosity of the suspension, such as sodium carboxymethyl cellulose, sorbitol, or dextran. Non-lipid polycationic amino polymers can also be used for delivery. Optionally, the suspension can also contain suitable stabilizers or agents that increase the solubility of the compounds to allow for the preparation of highly concentrated solutions.

Pharmaceutical compositions of the present invention can also be formulated to permit injectable, long-term, deposition.

5 The pharmaceutical compositions of the present invention can be administered topically.

A topical semi-solid ointment formulation typically contains a concentration of the active ingredient from about 1 to 20%, e.g., 5 to 10%, in a carrier such as a pharmaceutical cream base. Various formulations for topical use include 10 drops, tinctures, lotions, creams, solutions, and ointments containing the active ingredient and various supports and vehicles. In other transdermal formulations, typically in patch-delivered formulations, the pharmaceutically active compound is formulated with one or more skin penetrants, such 15 as 2-N-methyl-pyrrolidone (NMP) or Azone.

Inhalation formulations can also readily be formulated. For inhalation, various powder and liquid formulations can be prepared.

The pharmaceutically active compound in the 20 pharmaceutical compositions of the present invention can be provided as the salt of a variety of acids, including but not limited to hydrochloric, sulfuric, acetic, lactic, tartaric, malic, and succinic acid. Salts tend to be more soluble in aqueous or other protonic solvents than are the corresponding 25 free base forms.

After pharmaceutical compositions have been prepared, they are packaged in an appropriate container and labeled for treatment of an indicated condition.

The active compound will be present in an amount 30 effective to achieve the intended purpose. The determination of an effective dose is well within the capability of those skilled in the art.

A "therapeutically effective dose" refers to that amount of active ingredient - for example RGL3 protein, fusion protein, or fragments thereof, antibodies specific for RGL3, agonists, antagonists or inhibitors of RGL3 - which ameliorates 5 the signs or symptoms of the disease or prevents progression thereof; as would be understood in the medical arts, cure, although desired, is not required.

The therapeutically effective dose of the pharmaceutical agents of the present invention can be estimated 10 initially by *in vitro* tests, such as cell culture assays, followed by assay in model animals, usually mice, rats, rabbits, dogs, or pigs. The animal model can also be used to determine an initial useful concentration range and route of administration.

15 For example, the ED₅₀ (the dose therapeutically effective in 50% of the population) and LD₅₀ (the dose lethal to 50% of the population) can be determined in one or more cell culture of animal model systems. The dose ratio of toxic to therapeutic effects is the therapeutic index, which can be 20 expressed as LD₅₀/ED₅₀. Pharmaceutical compositions that exhibit large therapeutic indices are particularly useful.

The data obtained from cell culture assays and animal studies is used in formulating an initial dosage range for 25 human use, and preferably provides a range of circulating concentrations that includes the ED₅₀ with little or no toxicity. After administration, or between successive administrations, the circulating concentration of active agent varies within this range depending upon pharmacokinetic factors well known in the art, such as the dosage form employed, 30 sensitivity of the patient, and the route of administration.

The exact dosage will be determined by the practitioner, in light of factors specific to the subject requiring treatment. Factors that can be taken into account by

the practitioner include the severity of the disease state, general health of the subject, age, weight, gender of the subject, diet, time and frequency of administration, drug combination(s), reaction sensitivities, and tolerance/response to therapy. Long-acting pharmaceutical compositions can be administered every 3 to 4 days, every week, or once every two weeks depending on half-life and clearance rate of the particular formulation.

Normal dosage amounts may vary from 0.1 to 100,000 micrograms, up to a total dose of about 1 g, depending upon the route of administration. Where the therapeutic agent is a protein or antibody of the present invention, the therapeutic protein or antibody agent typically is administered at a daily dosage of 0.01 mg to 30 mg/kg of body weight of the patient (e.g., 1mg/kg to 5 mg/kg). The pharmaceutical formulation can be administered in multiple doses per day, if desired, to achieve the total desired daily dose.

Guidance as to particular dosages and methods of delivery is provided in the literature and generally available to practitioners in the art. Those skilled in the art will employ different formulations for nucleotides than for proteins or their inhibitors. Similarly, delivery of polynucleotides or polypeptides will be specific to particular cells, conditions, locations, etc.

Conventional methods, known to those of ordinary skill in the art of medicine, can be used to administer the pharmaceutical formulation(s) of the present invention to the patient. The pharmaceutical compositions of the present invention can be administered alone, or in combination with other therapeutic agents or interventions.

THERAPEUTIC METHODS

The present invention further provides methods of treating subjects having defects in RGL3 - e.g., in expression, 5 activity, distribution, localization, and/or solubility of RGL3 - which can manifest as a disorder of adrenal, adult liver, bone marrow, brain, fetal liver, heart, kidney, lung, placenta, colon, skeletal muscle and prostate function. As used herein, "treating" includes all medically-acceptable types 10 of therapeutic intervention, including palliation and prophylaxis (prevention) of disease.

In one embodiment of the therapeutic methods of the present invention, a therapeutically effective amount of a pharmaceutical composition comprising RGL3 protein, fusion, 15 fragment or derivative thereof is administered to a subject with a clinically-significant RGL3 defect.

Protein compositions are administered, for example, to complement a deficiency in native RGL3. In other embodiments, protein compositions are administered as a vaccine 20 to elicit a humoral and/or cellular immune response to RGL3. The immune response can be used to modulate activity of RGL3 or, depending on the immunogen, to immunize against aberrant or aberrantly expressed forms, such as mutant or inappropriately expressed isoforms. In yet other embodiments, protein fusions 25 having a toxic moiety are administered to ablate cells that aberrantly accumulate RGL3.

In another embodiment of the therapeutic methods of the present invention, a therapeutically effective amount of a pharmaceutical composition comprising nucleic acid of the 30 present invention is administered. The nucleic acid can be delivered in a vector that drives expression of RGL3 protein, fusion, or fragment thereof, or without such vector.

Nucleic acid compositions that can drive expression of RGL3 are administered, for example, to complement a deficiency in native RGL3, or as DNA vaccines. Expression vectors derived from virus, replication deficient retroviruses, 5 adenovirus, adeno-associated (AAV) virus, herpes virus, or vaccinia virus can be used - see, e.g., Cid-Arregui (ed.), Viral Vectors: Basic Science and Gene Therapy, Eaton Publishing Co., 2000 (ISBN: 188129935X) - as can plasmids .

Antisense nucleic acid compositions, or vectors that 10 drive expression of RGL3 antisense nucleic acids, are administered to downregulate transcription and/or translation of RGL3 in circumstances in which excessive production, or production of aberrant protein, is the pathophysiologic basis of disease.

15 Antisense compositions useful in therapy can have sequence that is complementary to coding or to noncoding regions of the RGL3 gene. For example, oligonucleotides derived from the transcription initiation site, e.g., between positions -10 and +10 from the start site, are particularly 20 useful.

Catalytic antisense compositions, such as ribozymes, that are capable of sequence-specific hybridization to RGL3 transcripts, are also useful in therapy. See, e.g., Phylactou, *Adv. Drug Deliv. Rev.* 44(2-3):97-108 (2000); Phylactou et al., 25 *Hum. Mol. Genet.* 7(10):1649-53 (1998); Rossi, *Ciba Found. Symp.* 209:195-204 (1997); and Sigurdsson et al., *Trends Biotechnol.* 13(8):286-9 (1995), the disclosures of which are incorporated herein by reference in their entireties.

Other nucleic acids useful in the therapeutic methods 30 of the present invention are those that are capable of triplex helix formation in or near the RGL3 genomic locus. Such triplexing oligonucleotides are able to inhibit transcription, Intody et al., *Nucleic Acids Res.* 28(21):4283-90 (2000);

McGuffie et al., *Cancer Res.* 60(14):3790-9 (2000), the disclosures of which are incorporated herein by reference, and pharmaceutical compositions comprising such triplex forming oligos (TFOs) are administered in circumstances in which 5 excessive production, or production of aberrant protein, is a pathophysiologic basis of disease.

In another embodiment of the therapeutic methods of the present invention, a therapeutically effective amount of a pharmaceutical composition comprising an antibody (including 10 fragment or derivative thereof) of the present invention is administered. As is well known, antibody compositions are administered, for example, to antagonize activity of RGL3, or to target therapeutic agents to sites of RGL3 presence and/or accumulation.

15 In another embodiment of the therapeutic methods of the present invention, a pharmaceutical composition comprising a non-antibody antagonist of RGL3 is administered. Antagonists of RGL3 can be produced using methods generally known in the art. In particular, purified RGL3 can be used to screen 20 libraries of pharmaceutical agents, often combinatorial libraries of small molecules, to identify those that specifically bind and antagonize at least one activity of RGL3.

In other embodiments a pharmaceutical composition comprising an agonist of RGL3 is administered. Agonists can be 25 identified using methods analogous to those used to identify antagonists.

In still other therapeutic methods of the present invention, pharmaceutical compositions comprising host cells that express RGL3, fusions, or fragments thereof can be 30 administered. In such cases, the cells are typically autologous, so as to circumvent xenogeneic or allotypic rejection, and are administered to complement defects in RGL3 production or activity.

In other embodiments, pharmaceutical compositions comprising the RGL3 proteins, nucleic acids, antibodies, antagonists, and agonists of the present invention can be administered in combination with other appropriate therapeutic agents. Selection of the appropriate agents for use in combination therapy can be made by one of ordinary skill in the art according to conventional pharmaceutical principles. The combination of therapeutic agents or approaches can act additively or synergistically to effect the treatment or prevention of the various disorders described above, providing greater therapeutic efficacy and/or permitting use of the pharmaceutical compositions of the present invention using lower dosages, reducing the potential for adverse side effects.

15

TRANSGENIC ANIMALS AND CELLS

In another aspect, the invention provides transgenic cells and non-human organisms comprising RGL3 isoform nucleic acids, and transgenic cells and non-human organisms with targeted disruption of the endogenous orthologue of the human RGL3 gene.

The cells can be embryonic stem cells or somatic cells. The transgenic non-human organisms can be chimeric, nonchimeric heterozygotes, and nonchimeric homozygotes.

25

DIAGNOSTIC METHODS

The nucleic acids of the present invention can be used as nucleic acid probes to assess the levels of RGL3 mRNA in tumor tissues and cells, and antibodies of the present invention can be used to assess the expression levels of RGL3 proteins in tumor tissues and cells to diagnose cancer.

The following examples are offered for purpose of illustration, not limitation.

EXAMPLE 1

5 Identification and Characterization of
 cDNAs Encoding RGL3 Proteins

Predicating our gene discovery efforts on use of genome-derived single exon probes and hybridization to genome-
10 derived single exon microarrays – an approach that we have previously demonstrated will readily identify novel genes that have proven refractory to mRNA-based identification efforts – we identified an exon in raw human genomic sequence that is particularly expressed in human adrenal, adult liver, bone
15 marrow, brain, fetal liver, heart, kidney, lung, placenta, and prostate.

Briefly, bioinformatic algorithms were applied to human genomic sequence data to identify putative exons. Each of the predicted exons was amplified from genomic DNA,
20 typically centering the putative coding sequence within a larger amplicon that included flanking noncoding sequence. These genome-derived single exon probes were arrayed on a support and expression of the bioinformatically predicted exons assessed through a series of simultaneous two-color
25 hybridizations to the genome-derived single exon microarrays.

The approach and procedures are further described in detail in Penn et al., "Mining the Human Genome using Microarrays of Open Reading Frames," *Nature Genetics* 26:315-318 (2000); commonly owned and copending U.S. patent application
30 nos. 09/864,761, filed May 23, 2001, 09/774,203, filed January 29, 2001, and 09/632,366, filed August 3, 2000, the disclosures of which are incorporated herein by reference in their entireties.

Using a graphical display particularly designed to facilitate computerized query of the resulting exon-specific expression data, as further described in commonly owned and copending U.S. patent application no. 09/864,761, filed May 23, 5 2001, 09/774,203, filed January 29, 2001 and 09/632,366, filed August 3, 2000, the disclosures of which are incorporated herein by reference in their entireties, one exon was identified that is expressed in all the human tissues tested.

Table 1 summarizes the microarray expression data 10 obtained using genome-derived single exon probe corresponding to exon five. Each probe was completely sequenced on both strands prior to its use on a genome-derived single exon microarray; sequencing confirmed the exact chemical structure of each probe. An added benefit of sequencing is that it 15 placed us in possession of a set of single base-incremented fragments of the sequenced nucleic acid, starting from the sequencing primer's 3' OH. (Since the single exon probes were first obtained by PCR amplification from genomic DNA, we were of course additionally in possession of an even larger set of 20 single base incremented fragments of each of the single exon probes, each fragment corresponding to an extension product from one of the two amplification primers.)

Signals and expression ratios are normalized values measured and calculated as further described in commonly owned 25 and copending U.S. patent application nos. 09/864,761, filed May 23, 2001, 09/774,203, filed January 29, 2001, 09/632,366, filed August 3, 2000, and U.S. provisional patent application no. 60/207,456, filed May 26, 2000, the disclosures of which are incorporated herein by reference in their entireties.

Table 1
Expression Analysis
Genome-Derived Single Exon Microarray

Amplicon 24488, Exon 5		
(TISSUE)	Signal	Ratio
Adrenal	4.83	-1.01
adult liver	4.42	-1.04
bone marrow	4.32	1.07
Brain	3.86	-1.01
fetal liver	3.84	-1.16
Heart	4.89	-1.08
Hela	3.68	-1.38
Kidney	5.24	1.03
Lung	4.76	-1.02
Placenta	4.86	-1.17
Prostate	5.22	-1.08

As shown in Table 1, significant expression of exon five was seen only in adrenal, adult liver, bone marrow, brain, fetal liver, heart, kidney, lung, placenta and prostate, as well as a cell line, HeLa. Adrenal, adult liver, bone marrow, brain, fetal liver, heart, kidney, lung, placenta and prostate specific expression was further confirmed by RT-PCR analysis (see below).

Marathon-Ready™ brain cDNA (Clontech Laboratories, Palo Alto, CA, USA) was used as a substrate for standard RACE (rapid amplification of cDNA ends) to obtain a cDNA clone that spans 2.5 kilobases and appears to contain the entire coding region of the gene to which the exon contributes; for reasons described below, we termed this cDNA RGL3. Oligonucleotides OL649 (5'-ATGGAGCGCACAGCAGGCAAAGAGCTGGCCCTG-3'; SEQ ID NO: 45) and OL654 (5'-CAGGCTGCCTGGGTGTTGTCCTCTTC-3'; SEQ ID NO: 46) were used to PCR out a 1236 bp fragment of the open reading frame (ORF) using protocols according to manufacturer's

instructions (Clontech). Using similar protocol, oligonucleotides OL761 (5'-TCAGCTTGGGGAGACAGACAGAGTGTCCGGGTCC-3'; SEQ ID NO: 47) and OL764 (5'-CTCTCCCAGGTCATTGAGCCACCAGCTGCCTCCTG-3'; SEQ ID NO: 48) were used to PCR out another 642 bp fragment of the open reading frame (ORF). Based on the sequences of these two fragments, OL650 (5'-CATCCTGTCCGCCCTGCAATCTAACCCCC-3'; SEQ ID NO: 49) and OL651 (5'-CCGATGAGAACAAACCACCTCAGCAGCAGAG-5'; SEQ ID NO: 50) were used to RACE out the 3' RGL3 sequence and OL697 (5'-CATTGGTGGCAGCAGAAAAGCCCAGCAGG-3'; SEQ ID NO: 51) and OL698 (5'-CAGCAGGCAGGCAGTGGGTACAAAGGTCC-3'; SEQ ID NO: 52) were used to RACE out the 5' RGL3 sequence.

The RGL3 cDNAs were sequenced on both strands using a MegaBACE™ 1000 sequencer (Amersham Biosciences, Sunnyvale, CA, USA). Sequencing both strands provided us with the exact chemical structure of the cDNA, which is shown in FIG. 3 and further presented in the SEQUENCE LISTING as SEQ ID NO: 1, and placed us in actual physical possession of the entire set of single-base incremented fragments of the sequenced clone, starting at the 5' and 3' termini.

As shown in FIG. 3, the RGL3 cDNA spans 2496 nucleotides and contains an open reading frame from nucleotide 25 through and including nt 2157 (inclusive of termination codon), predicting a protein of 710 amino acids with a (posttranslationally unmodified) molecular weight of 78.1 kD. The clone appears full length, with the reading frame opening starting with a methionine and terminating with a stop codon.

BLAST query of genomic sequence identified two BACs, spanning over 53 kb, that constitute the minimum set of clones encompassing the cDNA sequence. Based upon the known origin of the BACs (GenBank accession numbers AC008481.8 and AC024575.5), the RGL3 gene can be mapped to human chromosome 19p13.2.

Comparison of the cDNA and genomic sequences identified 19 exons. Exon organization is listed in Table 2.

Table 2**RGL3 Exon Structure**

Exon no.	cDNA range	genomic range	BAC accession
1	1-57	139344-139400	AC008481.8
2	58-171	139862-139975	
3	172-395	141587-141810	
4	396-449	141979-142032	
5	450-661	91429-91218	AC024575.5
6	662-804	82145-82003	
7	805-1020	81915-81700	
8	1021-1124	80707-80604	
9	1125-1209	80512-80428	
10	1210-1266	77988-77932	
11	1267-1353	77823-77737	
12	1354-1386	77529-77497	
13	1387-1508	77414-77293	
14	1509-1604	75579-75484	
15	1605-1673	75390-75322	
16	1674-1770	75233-75137	
17	1771-1923	72879-72727	
18	1924-2038	72640-72526	
19	2039-2496	69802-69345	

5

FIG. 2 schematizes the exon organization of the RGL3 clone.

At the top is shown the two bacterial artificial chromosomes (BACs), with GenBank accession numbers (AC008481.8 and AC024575.5), that span the RGL3 locus. The genome-derived 10 single-exon probe first used to demonstrate expression from

this locus is shown below the BACs and labeled "500". The 500 bp probe includes sequence drawn solely from exon five.

As shown in FIG. 2, RGL3 encodes a protein of 710 amino acids and is comprised of exons 1 - 19. The predicted 5 molecular weight of RGL3, prior to any post-translational modification, is 78.1 kD.

As further discussed in the examples herein, expression of RGL3 was assessed using hybridization to genome-derived single exon microarrays. Microarray analysis of exon 10 five showed expression in all tissues tested, including adrenal, adult liver, bone marrow, brain, fetal liver, heart, kidney, lung, placenta and prostate, as well as a cell line, HeLa. RT-PCR confirmed microarray data, and further provided expression data for colon and skeletal muscle.

15 The sequence of the RGL3 cDNA was used as a BLAST query into the GenBank nr and dbEst databases. The nr database includes all non-redundant GenBank coding sequence translations, sequences derived from the 3-dimensional structures in the Brookhaven Protein Data Bank (PDB), sequences 20 from SwissProt, sequences from the protein information resource (PIR), and sequences from protein research foundation (PRF). The dbEst (database of expressed sequence tags) includes ESTs, short, single pass read cDNA (mRNA) sequences, and cDNA sequences from differential display experiments and RACE 25 experiments.

BLAST search identified multiple human and mouse ESTs, two ESTs from pig (AW786464.1 and BF192995.1) and two ESTs from rat (BF566047.1 and H32403.1) as having sequence closely related to RGL3. BLAST also identified BC014426 as an 30 allelic variant of human RGL3, with a two base pair deletion of bases 992-993 of RGL3. BC014426 thus encodes a truncated form of RGL3, with only 352 amino acids.

Globally, the human RGL3 protein resembles mouse RGL3 and other RasGEF proteins. The closest homolog (ortholog) of RGL3 is mouse RGL3 (GenBank accession: AF237669.1), with 79 % amino acid identity and 86 % amino acid similarity over the 5 entire length of the two proteins.

Motif searches using Pfam (<http://pfam.wustl.edu>), SMART (<http://smart.embl-heidelberg.de>), and PROSITE pattern and profile databases (<http://www.expasy.ch/prosite>), identified several known domains shared with mouse RGL3 and 10 other RasGEF proteins.

FIG. 1 shows the domain structure of RGL3 and the alignment of the three domains with that of other RasGEF proteins.

As schematized in FIG. 1, the newly isolated gene 15 product shares certain protein domains and an overall structural organization with mouse RGL3 and other RasGEF proteins. The shared structural features strongly imply that RGL3 plays a role similar to that of mouse RGL3, as a guanine nucleotide exchange factor for the small GTPase Ral and a 20 downstream effector for both Rit and Ras. Through the mediation of cellular signaling, RGL3 likely regulates cellular proliferation and transformation.

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Like mouse RGL3 and other RasGEF molecules, RGL3 contains three functional domains. A RasGEFN domain is located

close to the N-terminal end (amino acid sequence 64 - 198), a RasGEF domain is found at the center of the molecule (amino acid sequence 243 - 506) and a RA (Ras association) domain is found at the C-terminal end (613 - 699).

5 <http://www.ncbi.nlm.nih.gov/Structure/cdd/wrpsb.cgi>. The RasGEFN domain is identified in a subset of guanine nucleotide exchange factor for Ras-like small GTPases. Recent crystal structure of the RasGEFN domain shows that this domain is alpha-helical and plays a "purely structural role". Boriack-
10 Sjodin et al, *Nature* 394, 337-343 (1998). The RasGEF domain may function as the prime site for guanine nucleotide binding and exchange and the RA is a putative binding site for RasGTP effectors. Shao et al., *J. Biol. Chem.* 275:26914-26924 (2000).

Other signatures of the newly isolated RGL3 protein
15 were identified by searching the PROSITE database,
<http://www.expasy.ch/tools/scnpsit1.html>. These include one N-glycosylation sites (339-342), three cAMP- and cGMP-dependent protein kinase phosphorylation site (374-377, 517-520, 523-526), fourteen protein kinase C phosphorylation sites (30-32, 20 36-38, 63-65, 99-101, 170-172, 277-279, 290-292, 342-344, 372-374, 388-390, 495-497, 512-514, 559-561, and 591-593), nine casein kinase II phosphorylation sites (40-43, 221-224, 247-250, 256-259, 277-280, 388-391, 402-405, 540-543, and 635-638), and thirteen N-myristoylation sites (24-29, 169-174, 181-186, 25 273-278, 283-288, 286-291, 303-308, 307-312, 410-415, 425-430, 554-559, 568-573, and 700-705).

Possession of the genomic sequence permitted search
for promoter and other control sequences for the RGL3 gene.
A putative transcriptional control region, inclusive of
30 promoter and downstream elements, was defined as 1 kb around
the transcription start site, itself defined as the first
nucleotide of the RGL3 cDNA clone. The region, drawn from
sequence of BAC AC008481.8 , has the sequence given in SEQ ID

NO: 44, which lists 1000 nucleotides before the transcription start site.

Transcription factor binding sites were identified using a web based program (<http://motif.genome.ad.jp/>),
5 including binding sites for Lyf1 (49-57 and 365-373), CdxA (320-326, 611-617 and 623-629, with numbering according to SEQ ID NO: 29), amongst others.

We have thus identified a newly described human gene, RGL3, which shares certain protein domains and an overall
10 structural organization with mouse RGL3. The shared structural features strongly imply that the RGL3 protein plays a role similar to mouse RGL3, as a guanine nucleotide exchange factor for the small GTPase Ral and a downstream effector for both Rit and Ras, making the RGL3 proteins and nucleic acids clinically
15 useful diagnostic markers and potential therapeutic agents for cancer.

EXAMPLE 2
Preparation and Labeling of
20 Useful Fragments of RGL3

Useful fragments of RGL3 are produced by PCR, using standard techniques, or solid phase chemical synthesis using an automated nucleic acid synthesizer. Each fragment is
25 sequenced, confirming the exact chemical structure thereof.

The exact chemical structure of preferred fragments is provided in the attached SEQUENCE LISTING, the disclosure of which is incorporated herein by reference in its entirety. The following summary identifies the fragments whose structures are
30 more fully described in the SEQUENCE LISTING:

SEQ ID NO: 1 (nt, full length RGL3 cDNA)
SEQ ID NO: 2 (nt, cDNA ORF)
SEQ ID NO: 3 (aa, full length RGL3 protein)

SEQ ID NO: 4 (nt, (nt 964 - 1023) portion of RGL3)
SEQ ID NO: 5 (aa, (aa 314 - 333) CDS entirely within
portion of RGL3)
SEQ ID NOS: 6 - 24 (nt, exons 1 - 19 (from genomic
5 sequence))
SEQ ID NOS: 25 - 43 (nt, 500 bp genomic amplicon centered
about exons 1 - 19)
SEQ ID NO: 44 (nt, 1000 bp putative promoter)
SEQ ID NO: 45 (nt, primer OL649 for PCR cloning of RGL3)
10 SEQ ID NO: 46 (nt, primer OL654 for PCR cloning of RGL3)
SEQ ID NO: 47 (nt, primer OL761 for PCR cloning of RGL3
and expression analysis)
SEQ ID NO: 48 (nt, primer OL764 for PCR cloning of RGL3)
SEQ ID NO: 49 (nt, primer OL650 for PCR cloning of RGL3)
15 SEQ ID NO: 50 (nt, primer OL651 for PCR cloning of RGL3)
SEQ ID NO: 51 (nt, primer OL697 for PCR cloning of RGL3)
SEQ ID NO: 52 (nt, primer OL698 for PCR cloning of RGL3)
SEQ ID NO: 53 (nt, primer OL694 for expression analysis
of RGL3)

20

Upon confirmation of the exact structure, each of the above-described nucleic acids of confirmed structure is recognized to be immediately useful as a RGL3-specific probe.

For use as labeled nucleic acid probes, the above-25 described RGL3 nucleic acids are separately labeled by random priming. As is well known in the art of molecular biology, random priming places the investigator in possession of a near-complete set of labeled fragments of the template of varying length and varying starting nucleotide.

30 The labeled probes are used to identify the RGL3 gene on a Southern blot, and are used to measure expression of RGL3 mRNA on a northern blot and by RT-PCR, using standard techniques.

EXAMPLE 3
Expression Analysis of RGL3 by RT-PCR

5 To explore the potential function of RGL3, the expression of RGL3 in human tissues was examined by PCR using marathon-ready cDNAs (Clontech). Oligonucleotides OL694 (5'-CCCTGTCCCCAACCTTGGCACCTTCCTTACG-3'; SEQ ID NO: 53) and OL761 (SEQ ID NO: 47) were used to amplify an 877 bp fragment of the
10 RGL3 ORF from human cDNAs of bone marrow, brain, colon, heart, kidney, liver, placenta, skeletal muscle and Hela cells. The PCR conditions were according to a touchdown PCR procedure. The tubes containing the oligonucleotides, cDNA and Taq polymerase were first incubated at 94°C for 15 seconds followed by 70°C
15 for 3 minutes, cycle 5 times. The tubes were then incubated at 94°C for 15 seconds followed by 68°C for 3 minutes, cycle 5 times. Finally the tubes were incubated at 94°C for 15 seconds followed by 66°C for 3 minutes, cycle 25 times. The result of
20 the expression profile is shown in FIG. 4. The abundance of PCR products indicate that RGL3 is highly expressed in all tissues and Hela cells except in skeletal muscle, where its expression is lower.

EXAMPLE 4
25 Production of RGL3 Protein

The full length RGL3 cDNA clone is cloned into the mammalian expression vector pcDNA3.1/HISA (Invitrogen, Carlsbad, CA, USA), transfected into COS7 cells, transfectants
30 selected with G418, and protein expression in transfectants confirmed by detection of the anti-XpressTM epitope according to manufacturer's instructions. Protein is purified using immobilized metal affinity chromatography and vector-encoded protein sequence is then removed with enterokinase, per

manufacturer's instructions, followed by gel filtration and/or HPLC.

Following epitope tag removal, RGL3 protein is present at a concentration of at least 70%, measured on a weight basis with respect to total protein (i.e., w/w), and is free of acrylamide monomers, bis acrylamide monomers, polyacrylamide and ampholytes. Further HPLC purification provides RGL3 protein at a concentration of at least 95%, measured on a weight basis with respect to total protein (i.e., w/w).

EXAMPLE 5
Production of Anti- RGL3 Antibody

Purified proteins prepared as in Example 4 are conjugated to carrier proteins and used to prepare murine monoclonal antibodies by standard techniques. Initial screening with the unconjugated purified proteins, followed by competitive inhibition screening using peptide fragments of the RGL3, identifies monoclonal antibodies with specificity for RGL3.

EXAMPLE 6
25 Use of RGL3 Probes and Antibodies
for Diagnosis of Cancer

After informed consent is obtained, samples are drawn from cancer tissue or cells and tested for RGL3 mRNA levels by standard techniques and tested additionally for RGL3 protein levels using anti- RGL3 antibodies in a standard ELISA.

EXAMPLE 7

Use of RGL3 Nucleic Acids, Proteins,
and Antibodies in Therapy

5 Once over-expression of RGL3 is detected in patient, RGL3 antisense RNA or RGL3-specific antibody is introduced into disease cells to reduce the amount of the protein.

Once mutations of RGL3 have been detected in patients, normal RGL3 is reintroduced into the patient's
10 disease cells by introduction of expression vectors that drive RGL3 expression or by introducing RGL3 proteins into cells. Antibodies for the mutated forms of RGL3 are used to block the function of the abnormal forms of the protein.

15

EXAMPLE 8

RGL3 Disease Associations

Diseases that map to the RGL3 chromosomal region are
20 shown in Table 3. Mutations of RGL3 might lead to the disease listed below. Alternatively, mutations of RGL3 might lead to some other human disorder(s) as well.

Table 3: Diseases mapped to human chromosome 19p13.2 (RGL3 region).		
mim_num	disease	chromosomal location
600209	EXOSTOSES, MULTIPLE, TYPE III; EXT3	19p
602077	ECTRODACTYLY, ECTODERMAL DYSPLASIA, AND CLEFT LIP/PALATE SYNDROME 2; EEC2	Chr.19

25 All patents, patent publications, and other published references mentioned herein are hereby incorporated by reference in their entireties as if each had been individually and specifically incorporated by reference herein. While preferred illustrative embodiments of the present invention are
30 described, one skilled in the art will appreciate that the

present invention can be practiced by other than the described embodiments, which are presented for purposes of illustration only and not by way of limitation. The present invention is limited only by the claims that follow.